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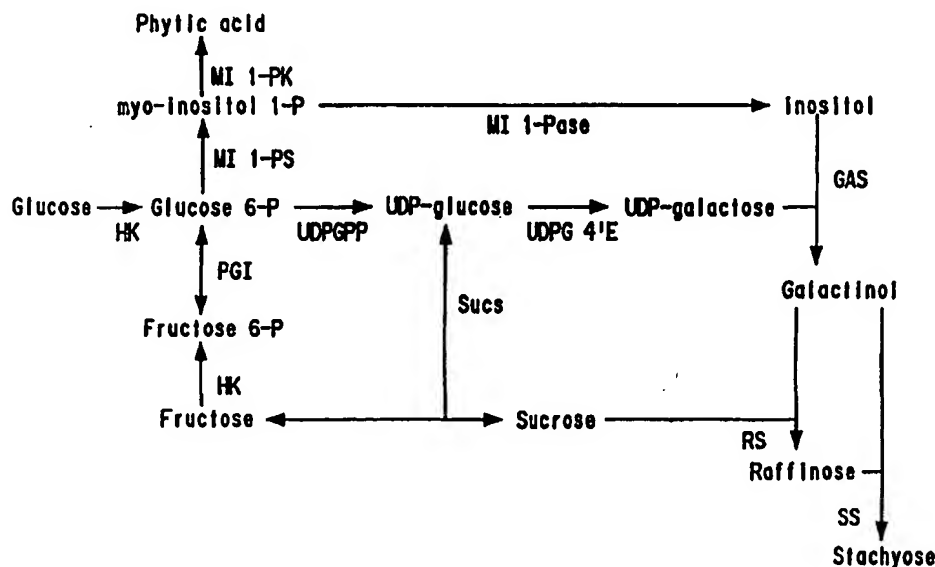
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(54) Title: SOYBEAN PLANT PRODUCING SEEDS WITH REDUCED LEVELS OF RAFFINOSE SACCHARIDES AND PHYTIC ACID



(57) Abstract

The instant invention pertains to the identity, characterization and manipulation of a soybean enzyme that results in the alteration of raffinose saccharide, sucrose, phytic acid and inorganic phosphate content of soybean seeds, thus leading to valuable and useful soybean products. The instant invention comprises soybean lines with decreased capacity for the synthesis of *myo*-inositol 1-phosphate in the tissue of developing seeds in comparison to seeds of other soybean lines. As taught herein, reduction of *myo*-inositol 1-phosphate synthase enzymatic activity by any of several means will result in soybean seeds displaying the instant phenotype.

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TITLESOYBEAN PLANT PRODUCING SEEDS WITH REDUCED LEVELS OF  
RAFFINOSE SACCHARIDES AND PHYTIC ACIDFIELD OF THE INVENTION

5           This invention is in the field of plant molecular biology and genetics. More specifically, this invention pertains to soybean protein and meal products having significantly lower contents of raffinose, stachyose and phytic acid and significantly higher contents of sucrose and inorganic phosphate as a function of the use of soybean lines having a heritable, decreased capacity for the production of *myo*-inositol  
10   l-phosphate in their seeds.

BACKGROUND OF THE INVENTION

          Raffinose saccharides are a group of D-galactose-containing oligosaccharides of sucrose that are widely distributed in plants. Raffinose saccharides are characterized by having the general formula  $O-\alpha\text{-D-galactopyranosyl}-(1\rightarrow6)_n-\alpha\text{-D-glucopyranosyl}-(1\rightarrow2)-\beta\text{-D-fructofuranoside}$  where  $n=1$  through  $n=4$  are known  
15   respectively as raffinose, stachyose, verbascose, and ajugose. In soybean seeds, raffinose and stachyose are the raffinose saccharides that are present in greatest quantity. In contrast, verbascose and ajugose are minor components and are generally not detected by standard analytical methods.

20           Extensive botanical surveys of the occurrence of raffinose saccharides have been reported in the scientific literature [Dey, P. M. In *Biochemistry of Storage Carbohydrates in Green Plants*, Academic Press, London, (1985) pp 53-129]. Raffinose saccharides are thought to be second only to sucrose among the nonstructural carbohydrates with respect to abundance in the plant kingdom. In fact,  
25   raffinose saccharides may be ubiquitous, at least among higher plants. Raffinose saccharides accumulate in significant quantities in the edible portion of many economically significant crop species. Examples include soybean (*Glycine max* L. Merrill), sugar beet (*Beta vulgaris*), cotton (*Gossypium hirsutum* L.), canola (*Brassica* sp.) and all of the major edible leguminous crops including beans (*Phaseolus* sp.),  
30   chick pea (*Cicer arietinum*), cowpea (*Vigna unguiculata*), mung bean (*Vigna radiata*), peas (*Pisum sativum*), lentil (*Lens culinaris*) and lupine (*Lupinus* sp.).

          Although abundant in many species, raffinose saccharides are an obstacle to the efficient utilization of some economically important crop species. Raffinose saccharides are not digested directly by animals, primarily because  $\alpha$ -galactosidase is  
35   not present in the intestinal mucosa [Gitzelmann and Auricchio. (1965) *Pediatrics* 36:231-236; Rutloff et al. (1967) *Nahrung*. 11:39-46]. However, microflora in the lower gut are readily able to ferment the raffinose saccharides which results in an acidification of the gut and production of carbon dioxide, methane and hydrogen [Murphy et al. (1972) *J. Agr. Food Chem.* 20:813-817; Cristofaro et al. In *Sugars in*

*Nutrition*, (1974) Chapter 20, 313-335; Reddy et al. (1980) *J. Food Science* 45:1161-1164]. The resulting flatulence can severely limit the use of leguminous plants in animal, including human, diets. It is unfortunate that the presence of raffinose saccharides restricts the use of soybeans in animal, including human, diets because otherwise this species is an excellent source of protein and fiber.

The soybean is well-adapted to machinery and facilities for harvesting, storing and processing that are widely available in many parts of the world. In the U.S. alone, approximately 28 million metric tons of meal were produced in 1988 [*Oil Crops Situation and Outlook Report*, Apr. 1989, U.S. Dept. of Agriculture, Economic Research Service]. Typically, hulls are removed and then the oil is extracted with hexane in one of several extraction systems. The remaining defatted flakes can then be used for a variety of commercial soy protein products [*Soy Protein Products, Characteristics, Nutritional Aspects and Utilization* (1987) Soy Protein Council]. Foremost among these in volume of use is soybean meal, the principle source of protein in diets used for animal feed, especially those for monogastric animals such as poultry and swine.

Although the soybean is an excellent source of vegetable protein, there are inefficiencies associated with its use that appear to be due to the presence of raffinose saccharides. Compared to maize, the other primary ingredient in animal diets, gross energy utilization for soybean meal is low [Potter and Potchanakorn. In: *Proceedings World Soybean Conference III*, (1984) 218-224]. For example, although soybean meal contains approximately 6% more gross energy than ground yellow corn, it has about 40 to 50% less metabolizable energy when fed to chickens. This inefficiency of gross energy utilization does not appear to be due to problems in digestion of the protein fraction of the meal, but rather due to the poor digestion of the carbohydrate portion of the meal. It has been reported that removal of raffinose saccharides from soybean meal by ethanol extraction results in a large increase in the metabolizable energy for broilers [Coon, C. N. et al. In: *Proceedings Soybean Utilization Alternatives*, University of Minnesota, (1988) 203-211]. Removal of the raffinose saccharides was associated with increased utilization of the cellulosic and hemicellulosic fractions of the soybean meal.

A variety of processed vegetable protein products are produced from soybean. These range from minimally processed, defatted items such as soybean meal, grits, and flours to more highly processed items such as soy protein concentrates and soy protein isolates. In other soy protein products the oil is not extracted, full-fat soy flour for example. In addition to these processed products, there are also a number of speciality products based on traditional Oriental processes, which utilize the entire bean as the starting material. Examples include soy milk, soy sauce, tofu, natto, miso, tempeh and yuba.

Examples of use of soy protein products in human foods include soy protein concentrates, soy protein isolates, textured soy protein, soy milk and infant formula. Facilities and methods to produce protein concentrates and isolates from soybeans are available across the world. One of the problems faced by producers of soy protein concentrates and isolates is the challenge of selectively purifying the protein away from the raffinose saccharides. Considerable equipment and operating costs are incurred as a result of removing the large amounts of raffinose saccharides that are present in soybeans.

The problems and costs associated with raffinose saccharides could be reduced or eliminated through the availability of genes that confer a reduction of raffinose saccharide content of soybean seeds. Such genes could be used to develop soybean varieties having inherently reduced raffinose saccharide content. Soybean varieties with inherently reduced raffinose saccharide content would improve the nutritional quality of derived soy protein products and reduce processing costs associated with the removal of raffinose saccharides. Low raffinose saccharide soybean varieties would be more valuable than conventional varieties for animal and human diets and would allow mankind to more fully utilize the desirable nutritional qualities of this edible legume.

*myo*-Inositol hexaphosphate (also known as "phytic acid") and raffinose saccharides share *myo*-inositol as a common intermediate in their synthesis [Ishitani, M. et al., (1996) *The Plant Journal* 9:537-548]. Like raffinose saccharides, phytic acid is a nearly ubiquitous component of angiosperm seeds [Raboy, V. In: *Inositol Metabolism in Plants* (1990) Wiley-Liss, New York, pp 55-76]. While phytic acid typically accounts for 50 to 70% of the total phosphate in seeds such as soybean and corn, that phosphate is only poorly available to mono-gastric animals. In addition to being only partially digestible, the presence of phytic acid in animal rations leads to excretion of other limiting nutrients such as essential amino acids, calcium and zinc [Mroz, Z. et al., (1994) *J. Animal Sci.* 72:126-132; Fox et al., In: *Nutritional Toxicology Vol. 3*, Academic Press, San Diego (1989) pp. 59-96]. Since soybean meal is a major portion of many animal feed rations, a meal with decreased amounts of phytic acid along with increased amounts of available phosphate should lead to improved feed efficiency in soy containing rations. Indeed enzymatic treatment of soybean meal containing rations to partially hydrolyze the phosphate groups from phytic acid improves both phosphate availability and the availability of other limiting nutrients [Mroz et al., *supra*; Pen et al., (1993) *Biotechnology* 11:811-814].

Surveys of commercial and wild soybean germplasm indicate that limited genetic variability for seed phytic acid content exists [Raboy et al., (1984) *Crop Science* 24:431-434]. In light of these factors, it is apparent that soybean plants with

heritable, substantially reduced levels of raffinose saccharides and phytic acid in their seeds are needed.

#### SUMMARY OF THE INVENTION

The instant invention pertains to a soybean plant with a heritable phenotype of  
5 (i) a seed phytic acid content of less than 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$ , the phenotype due to a decreased capacity for the synthesis of *myo*-inositol 1-phosphate in the seeds of the plant. The invention also pertains to seeds derived from this plant.

10 More specifically, this invention pertains to an isolated nucleic acid fragment encoding a soybean *myo*-inositol 1-phosphate synthase or its complement, and a chimeric gene comprising this nucleic acid fragment or a subfragment of this nucleic acid fragment, operably linked to suitable regulatory sequences, wherein expression of the chimeric gene results in a decrease in expression of a native gene encoding a  
15 soybean *myo*-inositol 1-phosphate synthase.

Another embodiment of the instant invention is an isolated nucleic acid fragment encoding a mutant *myo*-inositol 1-phosphate synthase having decreased capacity for the synthesis of *myo*-inositol-1-phosphate.

Yet another embodiment of the instant invention is a soybean plant having in  
20 its genome a chimeric gene comprising a nucleic acid fragment encoding a soybean *myo*-inositol 1-phosphate synthase or the complement of the nucleic acid fragment, operably linked to suitable regulatory sequences, wherein expression of the chimeric gene results in a decrease in expression of a native gene encoding a soybean *myo*-inositol-1-phosphate synthase.

25 Still another embodiment of the instant invention is a soybean plant homozygous for at least one gene encoding a mutant *myo*-inositol 1-phosphate synthase having decreased capacity for the synthesis of *myo*-inositol 1-phosphate, the gene conferring a heritable phenotype of (i) a seed phytic acid content of less than 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than  
30 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$ .

The instant invention also pertains to seeds and protein products derived from the seeds of any of the plants described above.

Yet another embodiment of the instant invention is a method for making a soybean plant with a heritable phenotype of (i) a seed phytic acid content less than  
35 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$ , the method comprising (a) crossing the soybean line LR33 or any soybean plant having in its genome a chimeric gene comprising a nucleic acid fragment encoding a soybean *myo*-inositol 1-phosphate synthase or the complement of the nucleic acid fragment,

operably linked to suitable regulatory sequences, or a soybean plant homozygous for at least one gene encoding a mutant *myo*-inositol 1-phosphate synthase having decreased capacity for the synthesis of *myo*-inositol 1-phosphate, with an elite soybean plant; and (b) selecting progeny plants of the cross of step (a) that have a heritable phenotype of (i) a seed phytic acid content less than 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$ .

The instant invention also embodies a method for producing a soy protein product comprising (a) crossing an agronomically elite soybean plant with LR33 or any of the soybean plants described above; (b) screening the seed of progeny plants obtained from step (a) for (i) a seed phytic acid content less than 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$ ; and (c) processing the seed selected in step (b) to obtain the desired soybean protein product.

Finally, the instant invention embodies a method of using a soybean plant homozygous for at least one gene encoding a mutant *myo*-inositol 1-phosphate synthase having decreased capacity for the synthesis of *myo*-inositol 1-phosphate, the gene conferring a heritable phenotype of (i) a seed phytic acid content less than 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$  to produce progeny lines, the method comprising (a) crossing a soybean plant comprising a mutant *myo*-inositol 1-phosphate synthase having decreased capacity for the synthesis of *myo*-inositol 1-phosphate with any soybean parent which does not comprise the mutation, to yield a F1 hybrid; (b) selfing the F1 hybrid for at least one generation; and (c) identifying the progeny of step (b) homozygous for at least one gene encoding a mutant *myo*-inositol 1-phosphate synthase having decreased capacity for the synthesis of *myo*-inositol 1-phosphate, the gene conferring a heritable phenotype of (i) a seed phytic acid content less than 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$ .

#### BRIEF DESCRIPTION OF THE DRAWINGS AND SEQUENCE LISTINGS

The invention can be more fully understood from the figure and the Sequence Listings which form a part of this application. The Sequence Listings contain the three letter codes for amino acids as defined in 37 C.F.R. 1.822 which are incorporated herein by reference.

Figure 1. Glucose metabolism to phytic acid, raffinose and stachyose in soybean seeds. The common co-factors, ATP, ADP, UTP, UDP, pyrophosphate and inorganic phosphate (Pi) are not shown. The enzymes are listed by abbreviation:

hexokinase (HK); phosphoglucosomerase (PGI); UDPglucose pyrophosphorylase (UDPGPP); UDPglucose 4' epimerase (UDPG 4'E); *myo*-inositol 1-phosphate synthase (MI 1-PS); *myo*-inositol 1-phosphatase (MI 1-Pase); *myo*-inositol 1-phosphate kinase (MI 1-PK); galactinol synthase (GAS); sucrose synthase (SucS); raffinose synthase (RS); and stachyose synthase (SS).

SEQ ID NO:1 is the 5' to 3' nucleotide sequence of the 1782 bases of the cDNA encoding the wild type soybean *myo*-inositol 1-phosphate synthase present in clone p5bmi-1ps.

SEQ ID NO:2 is the 510 amino acid sequence deduced from the open reading frame in SEQ ID NO:1.

SEQ ID NO:3 is the nucleotide sequence of the upstream (5') primer used in the isolation of the *myo*-inositol-1-phosphate synthase cDNA from soybean line LR33.

SEQ ID NO:4 is the nucleotide sequence of the downstream (3') primer used in the isolation of the *myo*-inositol 1-phosphate synthase cDNA from soybean line LR33.

SEQ ID NO:5 is the nucleotide sequence of the 1533 bases of cDNA encoding the LR33 *myo*-inositol 1-phosphate synthase present in clone LR33-10.

SEQ ID NO:6 is the 510 amino acid sequence deduced from the open reading frame in SEQ ID NO:5.

SEQ ID NO:7 is the nucleotide sequence of the upstream (5') primer used for PCR amplification of the wild type allele encoding *myo*-inositol 1-phosphate synthase from genomic DNA samples.

SEQ ID NO:8 is the nucleotide sequence of the upstream (5') primer used for PCR amplification of the LR33 allele encoding *myo*-inositol-1-phosphate synthase from genomic DNA samples.

#### BIOLOGICAL DEPOSITS

The following biological materials have been deposited under the terms of the Budapest Treaty at American Type Culture Collection (ATCC), 12301 Parklawn Drive, Rockville, MD 20852, and bear the following accession numbers:

Designation	Material	Accession Number	Date of Deposit
LR33	Seed	ATCC 97988	April 17, 1997
4E76	Seed	ATCC 97971	April 4, 1997
p5bmi-1ps	Plasmid	ATCC 97970	April 4, 1997

#### DETAILED DESCRIPTION

In the context of this disclosure, a number of terms shall be utilized. As used herein, "soybean" refers to the species *Glycine max*, *Glycine soja*, or any species that



is sexually cross compatible with *Glycine max*. A "line" is a group of plants of similar parentage that display little or no genetic variation between individuals for a least one trait. Such lines may be created by one or more generations of self-pollination and selection, or vegetative propagation from a single parent including by tissue or cell culture techniques. "Mutation" refers to a detectable and heritable genetic change (either spontaneous or induced) not caused by segregation or genetic recombination. "Mutant" refers to an individual, or lineage of individuals, possessing a mutation. A "population" is any group of individuals that share a common gene pool. In the instant invention, this includes M1, M2, M3, M4, F1, and F2 populations. As used herein, an "M1 population" is the progeny of seeds (and resultant plants) that have been exposed to a mutagenic agent, while "M2 population" is the progeny of self-pollinated M1 plants, "M3 population" is the progeny of self-pollinated M2 plants, and "M4 population" is the progeny of self-pollinated M3 plants. As used herein, an "F1 population" is the progeny resulting from cross pollinating one line with another line. The format used herein to depict such a cross pollination is "female parent\*male parent". An "F2 population" is the progeny of the self-pollinated F1 plants. An "F2-derived line" or "F2 line" is a line resulting from the self-pollination of an individual F2 plant. An F2-derived line can be propagated through subsequent generations (F3, F4, F5 etc.) by repeated self-pollination and bulking of seed from plants of said F2-derived line.

The term "nucleic acid" refers to a large molecule which can be single-stranded or double-stranded, composed of monomers (nucleotides) containing a sugar, a phosphate and either a purine or pyrimidine. A "nucleic acid fragment" is a fraction of a given nucleic acid molecule. A "subfragment" refers to a contiguous portion of a nucleic acid fragment comprising less than the entire nucleic acid fragment. "Complementary" refers to the specific pairing of purine and pyrimidine bases that comprise nucleic acids: adenine pairs with thymine and guanine pairs with cytosine. Thus, the "complement" of a first nucleic acid fragment refers to a second nucleic acid fragment whose sequence of nucleotides is complementary to the first nucleic acid sequence.

In higher plants, deoxyribonucleic acid (DNA) is the genetic material while ribonucleic acid (RNA) is involved in the transfer of the information in DNA into proteins. A "genome" is the entire body of genetic material contained in each cell of an organism. The term "nucleotide sequence" refers to the sequence of DNA or RNA polymers, which can be single- or double-stranded, optionally containing synthetic, non-natural or altered nucleotide bases capable of incorporation into DNA or RNA polymers. The term "oligomer" refers to short nucleotide sequences, usually up to 100 bases long. As used herein, the term "homologous to" refers to the relatedness between the nucleotide sequence of two nucleic acid molecules or between the amino

acid sequences of two protein molecules. Estimates of such homology are provided by either DNA-DNA or DNA-RNA hybridization under conditions of stringency as is well understood by those skilled in the art (Hames and Higgins, Eds. (1985) *Nucleic Acid Hybridisation*, IRL Press, Oxford, U.K.); or by the comparison of sequence similarity between two nucleic acids or proteins, such as by the method of Needleman et al. (*J. Mol. Biol.* (1970) 48:443-453). As used herein, "substantially similar" refers to nucleotide sequences that have more than 90% overall identity at the nucleotide level with the coding region of the claimed sequence, such as genes and pseudo-genes corresponding to the coding regions. The nucleic acid fragments described herein include molecules which comprise possible variations, both man-made and natural, such as but not limited to (a) those that involve base changes that do not cause a change in an encoded amino acid, or (b) which involve base changes that alter an amino acid but do not affect the functional properties of the protein encoded by the DNA sequence, (c) those derived from deletions, rearrangements, amplifications, random or controlled mutagenesis of the nucleic acid fragment, and (d) even occasional nucleotide sequencing errors.

"Gene" refers to a nucleic acid fragment that expresses a specific protein, including regulatory sequences preceding (5' non-coding) and following (3' non-coding) the coding region. "Native" gene refers to an isolated gene with its own regulatory sequences as found in nature. "Chimeric gene" refers to a gene that comprises heterogeneous regulatory and coding sequences not found in nature. "Endogenous" gene refers to the native gene normally found in its natural location in the genome and is not isolated. A "foreign" gene refers to a gene not normally found in the host organism but that is introduced by gene transfer. "Pseudo-gene" refers to a genomic nucleotide sequence that does not encode a functional enzyme.

"Coding sequence" refers to a DNA sequence that codes for a specific protein and excludes the non-coding sequences. It may constitute an "uninterrupted coding sequence", i.e., lacking an intron or it may include one or more introns bounded by appropriate splice junctions. An "intron" is a nucleotide sequence that is transcribed in the primary transcript but that is removed through cleavage and re-ligation of the RNA within the cell to create the mature mRNA that can be translated into a protein.

"Initiation codon" and "termination codon" refer to a unit of three adjacent nucleotides in a coding sequence that specifies initiation and chain termination, respectively, of protein synthesis (mRNA translation). "Open reading frame" refers to the coding sequence uninterrupted by introns between initiation and termination codons that encodes an amino acid sequence.

"RNA transcript" refers to the product resulting from RNA polymerase-catalyzed transcription of a DNA sequence. When the RNA transcript is a perfect copy of the DNA sequence, it is referred to as the primary transcript or it may be a

RNA sequence derived from posttranscriptional processing of the primary transcript and is then referred to as the mature RNA. "Messenger RNA (mRNA)" refers to the RNA that is without introns and that can be translated into protein by the cell. "cDNA" refers to a double-stranded DNA that is derived from mRNA. "Sense" RNA refers to an RNA transcript that includes the mRNA. "Antisense RNA" refers to an RNA transcript that is complementary to all or part of a target primary transcript or mRNA and that blocks the expression of a target gene by interfering with the processing, transport and/or translation of its primary transcript or mRNA. The complementarity of an antisense RNA may be with any part of the specific gene transcript, i.e., at the 5' non-coding sequence, 3' non-coding sequence, introns, or the coding sequence. In addition, as used herein, antisense RNA may contain regions of ribozyme sequences that increase the efficacy of antisense RNA to block gene expression. "Ribozyme" refers to a catalytic RNA and includes sequence-specific endoribonucleases.

As used herein, "suitable regulatory sequences" refer to nucleotide sequences in native or chimeric genes that are located upstream (5'), within, and/or downstream (3') to the nucleic acid fragments of the invention, which control the expression of the nucleic acid fragments of the invention.

"Promoter" refers to a DNA sequence in a gene, usually upstream (5') to its coding sequence, which controls the expression of the coding sequence by providing the recognition for RNA polymerase and other factors required for proper transcription. In artificial DNA constructs promoters can also be used to transcribe antisense RNA. Promoters may also contain DNA sequences that are involved in the binding of protein factors which control the effectiveness of transcription initiation in response to physiological or developmental conditions. It may also contain enhancer elements. An "enhancer" is a DNA sequence which can stimulate promoter activity. It may be an innate element of the promoter or a heterologous element inserted to enhance the level and/or tissue-specificity of a promoter. "Constitutive promoters" refers to those that direct gene expression in all tissues and at all times. "Tissue-specific" or "development-specific" promoters as referred to herein are those that direct gene expression almost exclusively in specific tissues, such as leaves or seeds, or at specific development stages in a tissue, such as in early or late embryogenesis, respectively. The term "expression", as used herein, refers to the transcription and stable accumulation of the sense (mRNA) or the antisense RNA derived from the nucleic acid fragment(s) of the invention that, in conjunction with the protein apparatus of the cell, results in altered levels of *myo-inositol 1-phosphate* synthase. "Antisense inhibition" refers to the production of antisense RNA transcripts capable of preventing the expression of the target protein. "Overexpression" refers to the production of a gene product in transgenic organisms that exceeds levels of

production in normal or non-transformed organisms. "Cosuppression" refers to the expression of a foreign gene which has substantial homology to an endogenous gene resulting in the suppression of expression of both the foreign and the endogenous gene. "Altered levels" refers to the production of gene product(s) in transgenic organisms in amounts or proportions that differ from that of normal or non-transformed organisms. The present invention also relates to vectors which include nucleotide sequences of the present invention, host cells which are genetically engineered with vectors of the invention and the production of polypeptides of the invention by recombinant techniques. "Transformation" herein refers to the transfer of a foreign gene into the genome of a host organism and its genetically stable inheritance. "Fertile" refers to plants that are able to propagate sexually.

"Raffinose saccharides" refers to the family of oligosaccharides with the general formula  $O-\alpha\text{-D-galactopyranosyl-(1}\rightarrow\text{6)}_n-\alpha\text{-D-glucopyranosyl-(1}\rightarrow\text{2)}-\beta\text{-D-fructofuranoside}$  where  $n=1$  to 4. In soybean seeds, the term refers more specifically to the members of the family containing one (raffinose) and two (stachyose) galactose residues. Although higher galactose polymers are known (e.g., verbascose and ajugose), the content of these higher polymers in soybean is below standard methods of detection and therefore do not contribute significantly to total raffinose saccharide content.

"Plants" refer to photosynthetic organisms, both eukaryotic and prokaryotic, whereas the term "Higher plants" refers to eukaryotic plants.

This invention provides a mutated form of a soybean gene and methods to improve the carbohydrate and phytic acid composition of soybean seeds and derived products. The invention teaches examples of a method to identify mutations in this gene and to use derived mutant soybean lines to reduce the raffinose saccharide and phytic acid content of soybean seeds. This invention also teaches methods of using gene silencing technology and the soybean gene sequence for *myo*-inositol-1-phosphate synthase discovered in the instant invention to reduce the raffinose saccharide content in soybean seeds.

Seeds derived from the plants of the present invention express an improved soluble carbohydrate content relative to commercial varieties. The improvements result in a reduced total raffinose plus stachyose content. The carbohydrate profile of these lines is dramatically different from the profiles seen in elite or germplasm lines used in or produced by other soybean breeding programs.

Two separate methods to produce the novel soybean genes of the present invention are taught. The first approach marks the first successful attempt to induce a mutation conferring low raffinose plus stachyose content. This approach resulted in the discovery of two major genes, one of which is described in detail in this invention, that can be used to develop soybean lines that are superior (in terms of reducing

combined raffinose and stachyose content) to any lines previously reported. The second approach utilizes the gene sequences taught in this invention applied in transgenic methods of specific gene silencing to achieve results similar to those obtained through random mutagenesis and screening.

5 Applicants initially sought to introduce random mutations in the genome of wild-type soybeans by chemical mutagenesis. The instant invention employed NMU (N-nitroso-N-methylurea) as the mutagenic agent, although other agents known to alter DNA structure and sequence could have been used. Following treatment NMU, soybean seeds were sown for several generation and screened for the desired  
10 phenotype; of primary importance was the alteration of raffinose saccharide content. Initial screening of mutagenized soybean populations revealed two lines, LR33 and LR28, that appeared to be low in raffinose saccharides (LR28 is disclosed in World Patent Publication WO93/07742). The low raffinose saccharide phenotype of LR33 was demonstrated to be inheritable by analysis of three subsequent generations of  
15 LR33 produced by self-fertilization (Table 1).

The physiological defect in LR33 leading to the unique phenotype displayed by this line was identified and characterized by conducting a series of elegant genetic and biochemical studies (see Example 2, *infra*). The defect in LR33 was shown to be genetically and biochemically distinct from the mutation in LR28 that leads to the low  
20 stachyose phenotype of that line. Moreover, the mutation in LR33 demonstrates greater pleiotrophy than the defect in LR28; the instant specification demonstrates that the LR33 phenotype includes not only reduced raffinose saccharide content, but also results in alterations in seed phytic acid, inorganic phosphate and sucrose levels. Further analyses confirmed that genetic information derived from LR33 alone could  
25 confer this unique phenotype on progeny soybean lines, and that the mutant gene or genes in LR33 are not simply genetic modifiers that enhance the phenotypic expression of genes derived from other mutant soybean lines.

The specific biochemical defect responsible for the heritable phenotype demonstrated by LR33 and its progeny has been identified. This was accomplished  
30 by consideration of the biosynthesis of raffinose saccharides and the control of phytic acid and inorganic phosphate levels in soybean seeds. Based upon these known biosynthetic pathways, a series of biochemical studies and subsequent molecular genetic analyses identified defect in LR33 seeds as an alteration in *myo*-inositol 1-phosphate synthase activity, leading to a decreased capacity for synthesis of  
35 *myo*-inositol 1-phosphate.

The biosynthesis of raffinose and stachyose has been fairly well characterized [see Dey, P. M. In Biochemistry of Storage Carbohydrates in Green Plants (1985)]. *Myo*-Inositol hexaphosphate or phytic acid and raffinose saccharides share *myo*-inositol as a common intermediate in their synthesis [Ishitani, M et al. The Plant

Journal (1996) 9:537-548]. Starting with glucose as a carbon source, the pathway describing the synthesis of phytic acid, raffinose and stachyose in maturing soybean seeds is shown in Figure 1.

By the interconversions shown in Figure 1, either glucose or sucrose can be the starting material for the polyol portion of phytic acid, all of the hexoses that make up raffinose and stachyose and the re-cycled portion of the galactose donor to raffinose synthase and stachyose synthase, *myo*-inositol. The end products of these interconversions that accumulate in mature, wild type soybean seeds are, in order of prominence by mass, sucrose, stachyose, phytic acid, and raffinose.

The committed reaction of raffinose saccharide biosynthesis involves the synthesis of galactinol (O- $\alpha$ -D-galactopyranosyl-(1 $\rightarrow$ 1)-*myo*-inositol) from UDPgalactose and *myo*-inositol. The enzyme that catalyzes this reaction is galactinol synthase. Synthesis of raffinose and higher homologues in the raffinose saccharide family from sucrose is catalyzed by the galactosyltransferases raffinose synthase and stachyose synthase.

Control over the ratio of these end products may be affected by altering the rate of conversion at many of the enzyme catalyzed steps in Figure 1. That control can be affected by altering enzyme expression level or by altering the intrinsic activity of the enzyme. The resulting mix of end products coming from the modified pathway may then comprise new proportions of the original end products as well as new product mixes which include accumulations of some of the normal intermediates. The exact mix and composition will depend upon both the enzyme which has been altered in its activity and the degree of that alteration.

The six enzymes *myo*-inositol 1-phosphate synthase, *myo*-inositol 1-phosphatase, UDP-glucose 4' epimerase, galactinol synthase, raffinose synthase, and stachyose synthase could be reduced in activity to decrease either raffinose or stachyose synthesis without decreasing sucrose content. Of these six, the three enzymes unique to raffinose and stachyose synthesis could be decreased in activity without decreasing phytic acid content. Only *myo*-inositol 1-phosphate synthase appears to be involved in the synthesis of all three end products and may therefore change the amount of all three end products simultaneously if its activity is decreased.

The instant invention teaches the ability to simultaneously reduce raffinose saccharide and phytic acid content and increase sucrose and inorganic phosphate content in soybean seeds by reducing *myo*-inositol 1-phosphate synthase activity in the cells of soybean seeds. The instant examples describe generation and discovery of a mutant form of this enzyme wherein a point mutation in the nucleotide sequence encoding this enzyme results in an amino acid substitution which, in turn, lowers intracellular enzymatic activity. It is well known to the skilled artisan that other mutations within the coding region for *myo*-inositol 1-phosphate synthase can result in

decreased enzymatic activity and thus result in the instant seed phenotype. Using well known techniques of heterologous gene expression and *in vitro* mutagenesis, and employing the various enzymatic assays described herein, the skilled artisan could identify other mutations within the *myo*-inositol 1-phosphate synthase coding region that result in decreased enzymatic activity without undue experimentation. These mutated *myo*-inositol 1-phosphate synthase genes could then be introduced into the soybean genome (see U.S. Pat. No. 5,501,967) and result in new soybean varieties displaying the instant phenotype.

Alternatively, gene silencing techniques such as antisense inhibition technology (U.S. Pat. No. 5,107,065) and cosuppression (U.S. Pat. No. 5,231,020) may be employed to reduce the intracellular *myo*-inositol 1-phosphate synthase activity in the cells of soybean seeds. The instant specification teaches the sequence of the gene encoding the wild type soybean *myo*-inositol 1-phosphate synthase enzyme. The skilled artisan will readily appreciate how to make and how to use chimeric genes comprising all or part of the wild type sequence or substantially similar sequences to reduce *myo*-inositol 1-phosphate synthase activity in soybean seeds.

Accordingly, the instant invention pertains to the identity, characterization and manipulation of a soybean enzyme that results in the alteration of raffinose saccharide, sucrose, phytic acid and inorganic phosphate content of soybean seeds, thus leading to valuable and useful soybean products. As taught herein, reduction of *myo*-inositol 1-phosphate synthase enzymatic activity by any of several means will result in soybean seeds displaying the instant phenotype.

#### EXAMPLES

The present invention is further defined in the following Examples, in which all parts and percentages are by weight and degrees are Celsius, unless otherwise stated. It should be understood that these Examples, while indicating preferred embodiments of the invention, are given by way of illustration only. From the above discussion and these Examples, one skilled in the art can ascertain, and without departing from the spirit and scope thereof, can make various changes and modifications of the invention to adapt it to various usages and conditions. Further, the present invention is not to be limited in scope by the biological materials deposited, since the deposited materials are intended to provide illustrations of materials from which many embodiments may be derived. All such modifications are intended to fall within the scope of the appended claims.

Usual techniques of molecular biology such as bacterial transformation, agarose gel electrophoresis of nucleic acids and polyacrylamide electrophoresis of proteins are referred to by the common terms describing them. Details of the practice of these techniques, well known to those skilled in the art, are described in detail in

[Sambrook, et al. (Molecular Cloning, A Laboratory Manual, 2nd ed. (1989), Cold Spring Harbor Laboratory Press]. Various solutions used in the experimental manipulations are referred to by their common names such as "SSC", "SSPE", "Denhardt's solution", etc. The composition of these solutions may be found by reference to Appendix B of Sambrook, et al. [*supra*].

#### EXAMPLE 1

### DISCOVERY OF A SOYBEAN GENE CONFERRING IMPROVED CARBOHYDRATE COMPOSITION

#### Assays for Raffinose Saccharide Content

The following assays were used to analyze soybean seeds for raffinose saccharide content. Prior to each of the analytical measures for determination of raffinose saccharide content, seeds were allowed to air dry for at least one week to a moisture content of approximately 8% and then stored at approximately 40 to 50°C and 40% relative humidity. Inventors' own measurements of many such air-dried samples indicated that the moisture content did not vary significantly from 8% (range of 7 to 10%) when stored at these conditions.

For each individual raffinose saccharide assay, typically five to ten soybeans from a given plant were ground in a Cyclotech 1093 Sample Mill (Tecator, Box 70 S-26301, Hoganas, Sweden) equipped with a 100 mesh screen to yield a seed powder that was then analyzed. In most cases, raffinose saccharide content was determined for seed or derived products containing an "as is" moisture content of approximately 8%. In these cases, "as is" values were converted to a dry basis (db) by dividing measurements by 0.92. For comparison among certain lines or among certain soy protein products, the ground seed powder was placed in a forced air oven at 45°C until the samples reached constant weight (0% moisture) prior to analysis. Hence, all raffinose saccharide measurements reported within this specification are on a common dry basis unless otherwise specified.

As describe below, three assays ("enzymatic", "TLC", and "HPLC") were used to determine raffinose saccharide content of soybean seeds. All three were performed on seed powder derived from the aforementioned grinding process.

In preparation for the "enzymatic" assay, after grinding each seed sample, approximately 30 mg of the resultant powder was weighed into a 13 x 100 mm screw cap tube and 1.6 mL of chloroform and 1.4 mL of methanol:water (4:3, v/v) was added. The precise powder weight of each sample was recorded and used to adjust the following assay results for sample to sample weight differences. The tubes were then capped, placed in racks and shaken on a rotary shaker for 60 min at 1800 rpm at room temperature. After extraction, the contents of the tubes were allowed to settle for 15 min. After settling, a 15 µL aliquot of the methanol:water phase was placed in a well of a 96 well microtiter plate and dried at 45°C for 20 min. The dried wells were



used as reaction vessels for the coupled "enzymatic" assay which employed  $\alpha$ -galactosidase and galactose dehydrogenase as described previously [Schiweck and Busching, (1969) *Zucker* 22:377-384; Schiweck and Busching, (1975) *Zucker* 28:242-243; Raffinose Detection Kit, Boehringer Mannheim GMBH, Catalog Number 428 167] with modifications of the assay conditions. The modifications of the assay included addition of Bovine Serum Albumin (15 mg/mL) to the assay and  $\alpha$ -galactosidase buffers, increasing the temperature and time of the  $\alpha$ -galactosidase incubation from room temperature to 45°C and 30 min, and increasing the time of the galactose dehydrogenase incubation from 20 min to 60 min, and using stachyose instead of raffinose for the  $\alpha$ -galactoside standard. After incubation, the absorbance at 340 nm of the samples was determined on a BIO-TEK® Model EL340 Microplate reader. The amount of  $\alpha$ -galactosides present in the samples was determined by comparison to known quantities of the stachyose standard. To facilitate the analysis of thousands of samples, enzymatic assays were replicated once. Lines that appeared to be low in raffinose saccharide content from the primary assay were subsequently reassayed in triplicate, beginning from the ground seed, if sufficient material was available. Lines whose composition was confirmed in the secondary assay were grown to maturity under field conditions and seed from the field-grown plants were assayed again. In cases where more specific information about the raffinose saccharide and galactinol profile was required, low raffinose saccharide lines identified by the enzymatic assay were reassayed using the HPLC assay (described below).

To facilitate the rapid selection of low raffinose saccharide germplasm, a thin layer chromatography "TLC" assay was developed. For this assay, about 60 mg of ground seed powder was placed into a 13 x 100 mm screw top test tube, to which 1 mL of 4:3 (v/v) methanol:water and 1 mL of chloroform were added. The tubes were capped, placed in racks and mixed on a rotary shaker for 60 min at 25 rpm at room temperature. After extraction the tubes were centrifuged at 2100 rpm for 5 min. A 4  $\mu$ L sample was taken from the methanol:water layer and placed on a 20 x 20 cm 'Baker' Silica Gel preadsorbent/channeled TLC plate with a 250 mm analytical layer. Samples were allowed to dry at room temperature and the plate was then placed in a TLC tank with a 3:4:4 (v/v/v) solution of Ethyl Acetate:Isopropanol:20% Acetic Acid (in water). The solution was allowed to soak up the analytical channels for 10 cm, at which time the plate was removed and allowed to air dry in a fume hood. The plate was then sprayed with an aniline-diphenylamine reagent to identify the carbohydrates in the sample. This reagent was prepared by mixing 1 mL of aniline with 100 mL of acetone, 1 gram of diphenylamine, and 10 mL of phosphoric acid. The plate was then placed in a 100°C oven for 15 min to dry and removed and allowed to cool down

before reading the analytical channels. Stachyose and raffinose content in the soybean samples were estimated by comparison to the elution pattern seen in pure standards.

A high performance anion exchange chromatography/pulsed amperometric assay, referred to herein as the "HPLC" assay, was used for determining the content of individual raffinose saccharides (e.g., stachyose and raffinose), galactinol, and for confirming the results of either the enzymatic or TLC assays. Conditions for the grinding and extraction of the seed were identical to those used for the previous "enzymatic" assay. A 750  $\mu$ L aliquot of the aqueous phase was removed and dried under reduced pressure at 80°C. The dried material was then dissolved in 2 mL of water and mixed vigorously for 30 sec. A 100  $\mu$ L aliquot was removed and diluted to 1 mL with water. The sample was mixed thoroughly again and then centrifuged for 3 min at 10,000 x g. Following centrifugation, a 20  $\mu$ L sample was analyzed on a Dionex™ PA1 column using 150 mM NaOH at 1.3 mL/min at room temperature. The Dionex™ PAD detector was used with E1 = 0.05 v, E2 = 0.60 v and E3 = -0.60 v and an output range of 3 mA. Galactinol, glucose, fructose, sucrose, raffinose, stachyose and verbascose were well separated by the chromatographic conditions. The carbohydrate content of the samples was determined by comparison to authentic standards.

Results obtained from the carbohydrate analyses were subjected to analysis of variance using the software SuperANOVA (Abacus Concepts, Inc., 1984 Bonita Avenue, Berkeley, CA 94704). When appropriate, Fisher's Protected LSD was used as the post-hoc test for comparison of means. In other comparisons, means were considered statistically significant if the ranges defined by their standard errors (SEM's) did not overlap. In cases where raffinose saccharide means were being compared to the mean of a control line, a mean was considered significantly lower than that of the control if said mean was at least three standard deviations below that of the control mean.

#### Mutagenesis and Selection of Mutants

Approximately 130,000 seeds (22.7 kg) of LR13 (a line essentially identical to Williams 82) were soaked in 150 L of tap water under continuous aeration for eight hours. Aeration was accomplished by pumping air through standard aquarium "airstones" placed in the bottom of the soaking vessel. Imbibed seeds were drained and transferred to 98 L of a 2.5 mM N-nitroso-N-methylurea (NMU) solution buffered at pH 5.5 with 0.1 M phosphate buffer under continuous aeration. Seeds remained in the NMU solution for three hours and were then put through a series of rinses to leach out the remaining NMU. For the first rinse, treated seeds were transferred to 45 L of tap water for 1 min. For the second rinse, seeds were transferred to 45 L of fresh tap water under continuous aeration for one hour. For the third rinse, seeds were transferred to 45 L of fresh tap water under continuous aeration for two hours. For the

fourth rinse, seeds were transferred to 45 L of fresh tap water under continuous aeration. One half of the seeds were removed from the fourth rinse after two hours (sub-population 1) while the other half of the seeds were removed from the fourth rinse after five hours (sub-population 2). After removal from the fourth rinse, seeds were drained of exogenous water and spread out on cardboard sheets to dry off in the sun for one hour. The imbibed M1 seeds were then field planted (Isabela, Puerto Rico, USA) in rows spaced 46 cm apart at a density of approximately 14 seeds per foot within the rows and a depth of 2.5 cm.

Two pools of M2 seeds (from sub-populations 1 and 2) were harvested in bulk from the M1 plants. Approximately 40,000 M2 seeds from sub-population 1 and 52,000 M2 seeds from sub-population 2 were planted at Isabela, Puerto Rico, USA. Within each sub-population, five pods from each of 3,000 M2 plants were harvested and bulked to obtain a bulk M3 seed population. M3 bulks were planted at Isabela, Puerto Rico. At maturity, seed from 5000 M3 plants were harvested individually to obtain 5000 M3:4 lines from each sub-population.

During the winter of 1991, a total of at least 8,000 M3:4 lines were screened to measure the content of raffinose saccharides using the enzymatic method described above. In the initial screening, two lines with decreased total raffinose saccharides were identified which proved heritable in a second, selfed generation. One of these lines, designated LR33, had reduced levels of both stachyose and raffinose in comparison to elite soybean cultivars grown as controls in the same environment. In comparison to the average values for three elite cultivars grown in the same environment, the stachyose content of LR33 was statistically significantly lower. The soluble carbohydrate content of bulked seeds harvested from selfed lines derived from LR33 for three generations is shown in Table 1.

TABLE 1  
Soluble carbohydrates in mature seeds of soybean lines derived by selfing line LR33

LINE	STACHYOSE	RAFFINOSE	GALACTINOL	SUCROSE
LR33	61	18	0	N.D. <sup>1</sup>
1ST-83	38	11	0	153
1991 Elite (avg.)	93	19	0	N.D. <sup>1</sup>
2ST-88	51	13	0	209
1992 Elite (avg.)	68	12	0	N.D. <sup>1</sup>
3ST-101	25	8	0	126
1993 Elite (avg.)	76	17	0	N.D. <sup>1</sup>

(All values are in  $\mu\text{moles g}^{-1}$ )

<sup>1</sup> N.D. Not determined

The subsequent lines (1ST-83, 2ST-88 and 3ST-101) were all derived by self pollination of LR33. Each of these lines had lower stachyose contents than did control lines.

### EXAMPLE 2

#### 5 IDENTIFICATION OF THE PHYSIOLOGICAL DEFECT RESPONSIBLE FOR THE LOW RAFFINOSE SACCHARIDE PHENOTYPE OF LR33

##### Genetic crosses using LR33 as a low raffinose saccharide parent

10 In an effort to further reduce the total raffinose saccharide content of soybean seeds, genetic crosses were made between LR33 and LR28. Line LR28 is described in World Patent Publication WO93/07742. Briefly, it is also a low combined raffinose plus stachyose line discovered in a screening procedure very similar to that described in Example 1. The low raffinose and stachyose content is consistent from one generation to the next. In addition, the galactinol content of the mature seeds of line LR28 and lines derived from it by crossing to other parents, is greatly elevated  
15 relative to wild type soybean seeds.

F1 plants from the cross of LR33 and LR28 were grown and self pollinated to produce segregating F2 progeny. Seeds of LR28-derived lines, elite cultivars and the segregating population resulting from the cross of LR28 and LR33 were planted in the same environment. F2:3 seeds were harvested from each F2 plant and screened for  $\alpha$ -galactoside content using the enzymatic assay described above. Low  $\alpha$ -galactoside  
20 selections from the preliminary screen were advanced to the HPLC assay to obtain complete raffinose saccharide profiles. Carbohydrate profiles from typical lines selected for low total  $\alpha$ -galactoside content are shown in Table 2.

**TABLE 2**

Soluble carbohydrates in mature seeds of soybean lines derived from crosses of line LR28 and LR33 and subsequent backcrosses to elite cultivars (LR33 and 5ST-1003 (an LR28 derived line) are included as examples of the cross parents; 2242 as an elite cultivar control.)

LINE	STACHYOSE	RAFFINOSE	GALACTINOL	SUCROSE
LR33	61	18	0	N.D. <sup>1</sup>
5ST-1441	57	18	0	N.D. <sup>1</sup>
5ST-1434	6	15	0	216
5ST-1309	0	5	0	287
2242	75	18	0	168
5ST-1003	19	4	65	200

25 (All values are in  $\mu\text{moles g}^{-1}$ )  
<sup>1</sup>N.D. Not determined

5ST-1441, 5ST-1434, and 5ST-1309 were all derived from the LR33 by LR28 cross. While line 5ST-1441 closely resembles the LR33 parent, surprisingly lines 5ST-1434 and 5ST-1309 are much lower in both raffinose and stachyose than either parental line and do not contain the elevated galactinol level characteristic of line LR28.

5 In vitro assay of activity of enzymes in the raffinose saccharide pathway

In studies designed to find the cause of the unexpected raffinose, stachyose and galactinol phenotype observed in some lines derived from the LR33 by LR28 cross, several enzyme activities and metabolite pools were measured in soybean seeds harvested just before the physiologically mature stage during the period of rapid  
10 raffinose saccharide biosynthesis. Seeds were removed from pods that had just begun to yellow. Seeds from such pods that were also losing green color or just beginning to yellow themselves were chosen for assay of the last three enzymes in raffinose saccharide biosynthesis (See Figure 1).

Seeds were removed from the pod, weighed to obtain fresh weight, then  
15 ground in a mortar and pestle in ten volumes of 50 mM HEPES-NaOH, pH 7 buffer which was also 5 mM in 2-mercaptoethanol. The ground samples were centrifuged at 10,000 x g for 10 min and the supernatant was desalted by passage through Sephadex G-25 which had been equilibrated in the grinding buffer.

For the assay of galactinol synthase, 10  $\mu$ L of the desalted extract was added  
20 to 90  $\mu$ L of the pH 7 HEPES buffer which was also 20 mM in *myo*-inositol, 10 mM in dithiothreitol, 1 mM in  $MnCl_2$  and 1 mM in UDP-[ $^{14}C$ ]galactose (1.25  $\mu$ Ci  $\mu$ mol $^{-1}$ ). The assay mixture was incubated for 10 min at 25°C then stopped by the addition of 400  $\mu$ L of ethanol. To the stopped reactions was added 200  $\mu$ L of Dowex AG-1x8 anion exchange resin (BioRAD) and the mixture was shaken for 25 min. The resin  
25 was removed by centrifugation and the supernatant was taken for scintillation counting. Non-anionic radioactivity was taken as a measure of galactinol synthesis.

For the assay of raffinose synthase and stachyose synthase, 50  $\mu$ L of the desalted extract was added to 50  $\mu$ L of 25 mM HEPES buffer at pH 7 that was also 10 mM in dithiothreitol, 10 mM in galactinol, and 40 mM in sucrose for the assay of  
30 raffinose synthase or 40 mM in raffinose for the assay of stachyose synthase. After a 1 h incubation at 25°C, the reactions were stopped by the addition of 40  $\mu$ L of ethanol and then placed in a boiling water bath for 1 min to precipitate proteins. The reaction mixes were centrifuged to clear, the supernatant was passed through a 0.22 micron filter and then reduced to dryness under vacuum. The residue was re-dissolved in  
35 0.5 mL of water and 20  $\mu$ L were separated on the Dionex HPLC system as described above for the quantitation of raffinose and stachyose. Results of an experiment done using seeds harvested from field grown plants in August of 1994 are given in Table 3.

TABLE 3

Activities of galactinol synthase, raffinose synthase, and stachyose synthase  
in yellowing seeds of three soybean lines  
(The wildtype control is line 1923. Enzyme activities are expressed as  $\mu$ mole  
of product produced per gram of fresh seed weight per hour under the assay conditions.  
Values are mean  $\pm$  standard deviation for five replicates.)

LINE	GALACTINOL SYNTHASE	RAFFINOSE SYNTHASE	STACHYOSE SYNTHASE
1923	7.5 $\pm$ 4.1	0.10 $\pm$ 0.05	0.65 $\pm$ 0.18
LR28	16.5 $\pm$ 5.2	0.004 $\pm$ 0.007	0.81 $\pm$ 0.24
LR28xLR33	22.6 $\pm$ 8.8	0.006 $\pm$ 0.008	0.87 $\pm$ 0.21

Of the three enzymes committed to raffinose and stachyose synthesis, only raffinose synthase shows a clear decrease in activity in the mutant, low raffinose saccharide lines in comparison to wild type. A mutation causing decreased activity of raffinose synthase is consistent with the position of that enzyme in the biosynthetic pathway and the accumulation of galactinol observed in lines which carry only LR28 as the source of the low raffinose saccharide phenotype. Galactinol and sucrose are the two substrates for raffinose synthase and both metabolites accumulate in LR28 containing lines (see Figure 1 and Table 2).

Despite the lower levels of stachyose, raffinose and galactinol obtained when LR33 lines were crossed with LR28 lines, the reduced raffinose synthase activity conferred by the LR28 mutation is the only altered activity. While decreased galactinol synthase activity seemed a likely candidate for the lesion in the LR33 line due the decreased galactinol content conferred by the mutation when in combination with LR28, there is no decrease in the measured, *in vitro* activity of that enzyme.

#### Assay of the free *myo*-inositol content of maturing soybean seeds

While there was no decrease in galactinol synthase *in vitro* in the LR33 by LR28 cross, the possibility that the *in vivo* activity of galactinol synthase in maturing seeds of LR33 derived lines is limited by the availability of one of its substrates was checked. The supply of UDP-galactose to galactinol synthase could be decreased by mutations in UDP-glucose 4' epimerase and the supply of *myo*-inositol could be limited by mutation effecting either its synthase or the specific phosphatase that produces the free inositol form (Figure 1). The *myo*-inositol pool was checked by assaying for the total free *myo*-inositol content in wild type and LR33 derived seeds. Three soybean lines, a wild type cultivar A2872, and two LR33xLR28-derived lines, 5ST-1434 and 5ST-1309, were grown in a growth room under 16 h days with a day/night temperature regime of 30°C/22°C. Seeds were taken from pods that had just begun to yellow and that were themselves turning from very light green to yellow. Three seeds from each line were ground in 8 mL of 80% methanol. The mixture was

centrifuged at 12,000 x g for 15 min and the supernatant decanted to a 50 mL flask. The extraction was repeated twice and the supernatants combined. The combined supernatants were reduced to dryness under vacuum at 40°C and the residues re-dissolved in 1 mL of water. The re-dissolved extracts were deionized by passage  
5 through 3 mL of mixed bed ion exchange resin (BioRad AG501x8) and the through flow plus 4 mL of wash was again reduced to dryness. The residue was re-dissolved in 500 µL of water and a 70 µL aliquot was separated by HPLC on a Zorbax Amino column (DuPont) eluted at 1 mL min<sup>-1</sup> with 70% acetonitrile. Sugars in the column eluate were detected by differential refractive index. Separation of standard mixtures  
10 of sucrose and *myo*-inositol showed that inositol emerges later and only partially separated from sucrose. For analysis of the seed extracts, 1 mL of eluate following the sucrose peak was trapped for re-injection on the Dionex PAD system described in Example 1. In the Dionex system, *myo*-inositol emerges well before sucrose and could be cleanly separated from sucrose contaminating the Zorbax Amino column  
15 fraction. By comparison to external standards of *myo*-inositol, the 5 µL injection from the wild type 2872 line contained 3.2 nmoles of *myo*-inositol while 5 µL injections from 5ST-1434 and 5ST-1309 contained 0.39 and 0.23 nmoles respectively.

A second experiment was performed to further characterize the difference in free *myo*-inositol content of wild type and LR33 derived seeds. Seven single seeds  
20 from a second control line (2242) and from the LR33 derived line 5ST-1434 were harvested according to the maturity criteria described above. The seeds were individually weighed, placed in 1.5 mL microfuge tubes, crushed with a spatula, frozen on dry ice and lyophilized. The dry residue was weighed and 1 mL of 80% methanol which contained 1 µmole of trehalose was added to act as a *myo*-inositol  
25 retention marker on the Zorbax amino column and as an internal standard. The extraction media was heated for 1 h at 60°C, re-ground with a small pestle in the microfuge tube and centrifuged to pellet the insoluble material. The supernatant was transferred to a second microfuge tube containing about 100 µL of mixed bed resin, the mixture was briefly shaken and 20 µL of the solution above the resin was removed  
30 and taken to dryness under vacuum. The residue was re-dissolved in 110 µL total volume and 55 µL were injected onto the Zorbax Amino column run as described above. The trehalose peak was trapped and 20 µL of the column fraction was re-injected on the Dionex system. The *myo*-inositol peak was quantitated by comparison to the trehalose internal standard. The mean ± standard deviation for the wild type  
35 *myo*-inositol content was 2.18±0.98 µmole (g dry wt)<sup>-1</sup> while the 5ST-1434 seeds contained 0.77±0.32 µmole (g dry wt)<sup>-1</sup>. Both experiments indicate that seeds which carry the LR33 mutation contain less *myo*-inositol during the period of rapid raffinose saccharide synthesis. In neither experiment was the total *myo*-inositol pool totally absent. Such experiments can only measure the total seed pool of metabolites

however and *myo*-inositol has other fates which could further limit its use by galactinol synthase (Figure 1).

The effect of *myo*-inositol perfusion on the *in vivo* labeling of raffinose saccharides

To ascertain whether or not the observed decrease in free *myo*-inositol content of the LR33 derived seeds might be the cause of the decreased raffinose saccharide content at maturity, tissue slice feeding studies were performed. Four seeds each from wild type line 2242 and LR33-derived line 5ST-1434 were harvested by the maturity criterion described above. The seed coats were removed and the cotyledons rinsed in 5 mM potassium phosphate buffer at pH 5.5 then sliced into approximately 1 mm slices and again rinsed with buffer. The tissue slices were divided into two groups. One group was immersed in potassium phosphate buffer and the second group was immersed in the potassium phosphate buffer containing 50 mM *myo*-inositol. The tissue slices were vacuum-infiltrated and incubated for 30 min at room temperature. After the pre-incubation period, 5  $\mu$ Ci of  $^{14}$ C-glucose was added to each grouping and the tissue was again vacuum-infiltrated. Ten tissue slices from each group were taken at 2 h and at 8 h after addition of the labeled glucose. The tissue slices were placed in tarred 1.5 mL microfuge tube to obtain fresh weight and ground in 300  $\mu$ L of 80% methanol. The tubes were centrifuged, the supernatant removed to a second tube, the extraction was repeated twice more and the supernatants were combined. The combined supernatants were reduced to dryness under vacuum, re-dissolved in 50% acetonitrile and separated on the Zorbax amino HPLC system. Fractions were collected for scintillation counting at 15 second intervals through the region of the chromatogram containing glucose, sucrose, raffinose and galactinol and at 1 min intervals through the region containing stachyose. Radioactive fractions corresponding to the sugar standard peaks were grouped for analysis. The results are expressed as per cent of the radiolabel in the four product sugars in Table 4.

Table 4

Radiolabel in sucrose, galactinol, raffinose and stachyose expressed as per cent of label in those sugars combined for control and an LR33 derived line with and without pre-incubation with *myo*-inositol

	WILD TYPE TISSUE SLICES				5ST-1434 TISSUE SLICES			
	2 h	2 h +inos	8 h	8 h +inos	2 h	2 h +inos	8 h	8 h +inos
SUCROSE	52.4	33.4	41.0	23.8	86.8	43.5	79.8	21.8
GALACTINOL	13.4	24.8	8.0	11.1	3.2	27.6	1.5	9.5
RAFFINOSE	19.8	20.7	17.3	17.7	4.5	15.2	3.1	13.0
STACHYOSE	14.4	21.0	33.6	47.4	5.4	13.7	15.9	55.6



Both wild type and 5ST-1434 tissue slices convert label from supplied  $^{14}\text{C}$ -glucose to sucrose and then into galactinol, raffinose, and stachyose. Without the addition of exogenous *myo*-inositol, the line containing the LR33 mutation (5ST-1434) converts very little label into the raffinose saccharides (20.5% after 8 h) in comparison to the wild type line (58.9% after 8 h). Both genotypes respond to exogenous *myo*-inositol by converting more of the supplied label to raffinose saccharide sugars. With the addition of *myo*-inositol the two genotypes become essentially equal in their ability to convert the supplied label first into galactinol and then into raffinose and stachyose. The rate of conversion is increased even in the wild type line in comparison to the non-infused control.

It appears that the supply of *myo*-inositol to galactinol synthase is always a limitation to the rate at which raffinose and stachyose can be synthesized. The presence of the LR33 mutation makes that limitation much greater.

The labeling pattern is surprising in that there is no buildup of label in galactinol as would be expected if the LR28 mutation in raffinose synthase were still effective in slowing flux through that step in the combined mutant lines. While the addition of *myo*-inositol can be expected to increase flux through galactinol synthase to galactinol, the label should have accumulated there due to the decreased activity in raffinose synthase. The absence of that accumulation calls into question either the effectiveness or the presence of the LR28 mutation in this line.

The influence of the LR33 mutation on seed phytic acid and inorganic phosphate levels

The above results suggest that the LR33 mutation resides before free *myo*-inositol in the pathway shown in Figure 1. To further limit the possible site of the mutation, other metabolites in that portion of the pathway were measured.

Since soybean seeds contain phytic acid at levels of 20 to 30  $\mu\text{moles g}^{-1}$  at maturity and since 50 to 70% of the total seed phosphate is contained in phytic acid [Raboy et al. (1984) *Crop Science* 24:431-434], it seemed likely that mutations effecting *myo*-inositol synthesis might also effect seed phytic acid and inorganic phosphate levels. The levels of phytic acid and inorganic phosphate in three wild type soybean lines and three lines that contain the LR33 mutation were assayed in dry seeds by a modification of the method described by Raboy [*supra*]. Approximately twenty seeds from each line were ground in a small impact mill and 100 mg of the resulting powder was weighed into a 15 mL screw cap tube. Five mL of 0.4 N HCl with 0.7M  $\text{Na}_2\text{SO}_4$  was added to the powder and the mixture was shaken overnight on a rocker platform. The tubes were centrifuged at 3900 x g for 15 min and 2 mL of the supernatant was removed to second, glass tube. Two mL of water and 1 mL of 15 mM  $\text{FeCl}_2$  were added and the tubes were heated for 20 min in a boiling water bath. The tubes were centrifuged as above to precipitate the iron:phytic acid complex

and the supernatant was discarded. The pellets were resuspended in 2 mL of 0.2 N HCl and heated for 10 min in the boiling water bath. The precipitate was again removed by centrifugation and re-suspended in 2 mL of 5 mM EDTA.

Fifteen  $\mu$ L of the EDTA suspension was taken for wet ashing to obtain phytic acid phosphate. To each tube containing the 15  $\mu$ L aliquot was added 10  $\mu$ L of a 10% solution of calcium nitrate. The water was allowed to evaporate and the residue was heated in a flame until a white ash remained in the tube. The ash was dissolved in 0.3 mL of 0.5 N HCL and heated at 90°C for 20 to 30 min. Acid molybdate reagent (0.7 mL of 0.36% ammonium molybdate and 1.42% ascorbic acid in 0.86 N sulfuric acid) was added and the color was allowed to develop for 1 h before reading at 820 nm.

Inorganic phosphate was determined by adding 0.3 mL of 0.5 N HCl and 0.7 mL of the phosphate color reagent to either 20 or 40  $\mu$ L aliquots of the initial 5 mL extract. Potassium phosphate standards were developed at the same time. The experiment was repeated three times and the results are given in Table 5.

Table 5

Seed phosphate in inorganic phosphate and in phytic acid expressed as  $\mu$ moles phosphate  $g^{-1}$   
(Values are mean  $\pm$  standard deviation where more than two replicates were obtained  
or the average value when two replicates were obtained.)

SEED LINE	GENOTYPE	REPLICATES	INORGANIC PHOSPHATE	PHYTIC ACID PHOSPHATE
2872	WILD TYPE	7	2.7 $\pm$ 3	125 $\pm$ 8.4
1923	WILD TYPE	1	2.0	126
1929	WILD TYPE	2	1.5	155
5ST-1309	LR33	4	76.0 $\pm$ 5.7	62.5 $\pm$ 12.4
GxE-117	LR33	7	36.7 $\pm$ 6.4	55.8 $\pm$ 13.9
GxE-76	LR33	6	32.2 $\pm$ 3.5	72.8 $\pm$ 14

The LR33 mutation also causes a decrease in seed phytic acid level (expressed as phosphate in the phytic acid fraction) of about two fold with a concomitant increase in the seed inorganic phosphate content of about 15 to 25 fold depending upon the genetic background in which the mutation is contained. Based on the pathway in Figure 1, the decreased phytic acid content, along with the decreased free *myo*-inositol content are most likely to be caused by a mutation decreasing the activity of *myo*-inositol 1-phosphate synthase.

The exact cause of the increase in the inorganic phosphate content is not known, however it has long been assumed that phytic acid functions as a phosphate storage form in seeds and other parts of the plant. It is possible that when the

preferred storage platform (*myo*-inositol) is unavailable, incoming inorganic phosphate has no other major fate and simply accumulates.

### EXAMPLE 3

#### IDENTIFICATION OF THE SEED PHENOTYPE OF THE HOMOZYGOUS LR33 LINE

Several additional soybean lines were analyzed for seed inorganic phosphate. One hundred mg of ground seed was extracted with 1 mL of hot water and centrifuged to remove the insoluble matter. The supernatant was extracted with 100  $\mu$ L of methylene chloride to remove lipid material and the remainder of the aqueous extract was made to 10% with trichloroacetic acid. The solution was centrifuged to remove proteins and 5  $\mu$ L of the supernatant was analyzed for inorganic phosphate as described in Example 2. Table 6 gives the results of those assays.

Table 6

The inorganic phosphate content ( $\mu$ mole g<sup>-1</sup>) of seeds from wild type, LR28, LR33 and LR28xLR33 soybean plants

LINE	GENOTYPE	INORGANIC PHOSPHATE
A2872	wildtype	12.7
LR28	LR28	12.7
LOW2	LR28	12.9
5ST-1190	LR28	14.4
5ST-1191	LR28	14.0
5ST-1441	LR33	23.4
5ST-1309	LR28xLR33	117.6
5ST-1310	LR28xLR33	113.9
5ST-1433	LR28xLR33	74.2
5ST-1434	LR28xLR33	57.1
GxE-117	LR28xLR33	74.3
GxE-305	LR28xLR33	70.7
GxE-76	LR28xLR33	79.1
GxE-77	LR28xLR33	58.4

All of the lines which contain the LR28 mutation alone have inorganic phosphate levels equal to the wild type controls. Line 5ST-1441 was derived from LR33 (Example 2 above) and has the moderate reduction in stachyose that was originally associated with the mutation. 5ST-1441 has only a slightly increased inorganic phosphate content. Despite the fact that neither parent is high in inorganic phosphate, all the lines derived from the cross of LR33 by LR28 have greatly elevated levels of inorganic phosphate.

The intermediate inorganic phosphate phenotype of line 5ST-1441 that contains only the LR33 mutation was unexpected. It may indicate that both the LR28 and the LR33 mutation must be present to produce the low phytic acid/high inorganic phosphate phenotype. If however the two mutations are in fact in the structural genes encoding the two activities that seem to be decreased, they are not related in the metabolic pathway in a manner that would suggest an additive effect on alteration of phytic acid synthesis. If the mutation in LR33 effects either free or total *myo*-inositol production, it alone should be responsible for the low phytic acid and resultant high inorganic phosphate phenotype.

An alternate explanation may be that the line 5ST-1441 is not homozygous for the LR33 mutation and that analysis of the bulk seed gives only the average phenotype for the wild type, homozygous mutant and heterozygous seeds. To check this possibility, thirteen single seeds from the bulk sample of 5ST-1441 seed were weighed, ground and extracted with hot water. Protein was not precipitated and inorganic phosphate was measured as before. The results are shown in Table 7.

Table 7

Approximate inorganic phosphate content ( $\mu\text{moles g}^{-1}$ ) for thirteen individual seeds from line 5ST-1441  
(Actual values are elevated in comparison to the results of Table 6 due to the presence of protein in the samples.)

SEED NUMBER	INORGANIC PHOSPHATE
5ST-1441-a	21.9
5ST-1441-b	30.4
5ST-1441-c	25.4
5ST-1441-d	81.1
5ST-1441-e	29.7
5ST-1441-f	25.7
5ST-1441-g	25.5
5ST-1441-h	30.8
5ST-1441-i	28.2
5ST-1441-j	123.5
5ST-1441-k	27.9
5ST-1441-l	115.9
5ST-1441-m	27.6

Seeds lettered d, j, and l have about three fold more inorganic phosphate than the remaining ten seeds analyzed. The ratio approximates that expected from a segregating population of seeds in which the mutant phenotype of high inorganic phosphate is recessive and due to the action of a single gene. To check this

- hypothesis, twenty seeds from the bulk sample of 5ST-1441 were imbibed in water at room temperature for 6 h. Excess water was blotted away and a sample of the cotyledons at the end away from the embryonic axis was cut off. The tissue pieces were weighed, placed in 1.5 mL microfuge tubes and ground in sufficient 70% methanol to give 10 mg fresh wt per 100  $\mu$ L of extracted volume. Phosphate in the supernatant present after centrifugation was measured as described above and the results are shown in Table 8.

Table 8

The inorganic phosphate content of 20 partial seeds from a segregating population of seeds from the line 5ST-1441  
(Results are expressed as  $\mu$ mole per g fresh weight ( $\mu$ mole gfw $^{-1}$ )).

SEED NO.	PO <sub>4</sub> ( $\mu$ mole gfw $^{-1}$ )
1	4.9
2	4.3
3	5.3
4	40.1
5	6.1
6	5.0
7	37.1
8	9.9
9	5.9
10	3.8
11	26.9
12	4.7
13	5.5
14	3.9
15	5.2
16	17.7
17	5.9
18	4.9
19	2.5
20	28.9

- Seeds 4, 7, 11, 16 and 20 were planted in pots in a growth room. Seeds 7, 11, 16, and 20 survived and were grown to maturity under the growth conditions described in Example 2. Bulk seeds from the mature plants and two wild type controls (A2872) were harvested after dry down and twenty seeds from each plant were ground in bulk for analysis of soluble carbohydrates and phytic acid using the

methods described in Examples 1 and 2. The total seed phytic acid content was calculated as the (phytic acid phosphate content)/6. The results are shown in Table 9.

Table 9

Soluble carbohydrate content and phytic acid content (expressed as  $\mu\text{mole g}^{-1}$ ) for four plants from a segregating population of seeds carrying the LR33 mutation and selected for elevated inorganic phosphate in partial seed analysis along with two control plants (Phytic acid values are the average of two replicates.)

LINE	PHYTIC ACID	STACHYOSE	RAFFINOSE	GALACTINOL	SUCROSE
A2872	32.3	71	19	3	166
A2872	30.8	67	24	0	144
LR33-7	23.7	40	16	0	155
LR33-11	8.8	4	13	0	212
LR33-16	9.5	4	13	2	209
LR33-20	7.6	4	11	0	208

- 5 While the soluble carbohydrate phenotype of the plant grown from seed number 7 (LR33-7) is very similar that ascribed to the LR33 mutation as shown in Table 1, the other plants have much lower levels of the raffinose saccharides and have much higher levels of sucrose. These levels of soluble carbohydrate are very similar to the phenotype ascribed to some plants in the mutant combination LR28xLR33 in
- 10 Table 2. The most probable explanation for this discrepancy is that the bulk seed analyzed to give the data in Table 1 came from plants segregating for the LR33 mutation rather than plants homozygous for the mutation. When plants homozygous for the mutation are selected, as was done in the case of plants 11, 16 and 20 in this example, the true, homozygous seed phenotype is observed. The previous assumption
- 15 that both the LR28 and the LR33 mutations needed to be present to produce the combination of low raffinose, stachyose and galactinol was incorrect. While lines homozygous for the LR33 mutation were not obtained from selfing the originally identified mutant line, they were obtained from segregants arising from the cross of that line with LR28. While some of the progeny of the cross likely do contain both
- 20 mutations, the phenotype from the LR33 mutation is dominant to that arising from the LR28 mutation. That dominance stems primarily from the location of the LR33 mutation upstream in the carbon flow from the LR28 mutation (see Figure 1 and Example 2).

- 25 The reason that the homozygous LR33 phenotype remained elusive may be due to germination problems encountered with the original mutant line. In bulk field segregation conditions the loss of 25% (that fraction that would have been homozygous for the mutation) of the planted seed from selfing LR33 could have been missed. Harvested plants from the bulk population would have then been depleted in

homozygotes but the mutant gene would be retained in the population in the heterozygous state. We surmise that out crossing to a genetic background more conducive to allowing LR33 homozygotes to emerge in field conditions allowed recovery of the homozygote. That the initial out crosses were to lines carrying  
5 another mutation in the raffinose saccharide biosynthetic pathway were incidental to the original intent of the cross and not essential either for the superior phenotype or the improved field emergence.

The LR33 mutation is thus capable of producing soybean plants that produce seeds with less than 5  $\mu$ moles of stachyose per gram of seed, less than 20  $\mu$ moles of  
10 raffinose per gram of seed, more than 200  $\mu$ moles of sucrose per gram of seed and less than 10  $\mu$ moles of phytic acid phosphate per gram of seed.

#### EXAMPLE 4

##### SOLUBLE CARBOHYDRATE AND PHYTIC ACID CONTENT OF A SOYBEAN LINE CONTAINING THE LR33 MUTATION

15 The low raffinose saccharide lines LR33 and LR28 were crossed and segregating F2 plants were selected for low levels of raffinose, stachyose and galactinol in the seed of the F2 plants by the HPLC assay described in Example 1. Selected lines derived from this cross were then crossed to elite cultivar parents and the progeny of those crosses were selected in the F2 generation by the same process.  
20 Among these lines containing low levels of raffinose saccharides, several were chosen for advancement based on the agronomic characteristics of the vegetative plant. One such line, designated 4E76, was subsequently test crossed to another elite soybean cultivar and single seeds from the selfed F1 plant derived from that cross were analyzed for soluble carbohydrates and for inorganic phosphate content. The seeds  
25 fell into two classes, those with very low inorganic phosphate and wild type levels of raffinose plus stachyose, and a smaller class of seeds with elevated levels of inorganic phosphate and very low levels of raffinose plus stachyose. The ratio of these two classes was very near the 3:1 ratio expected of the recessive LR33 mutation. There was no evidence of segregation of LR28 which is the other mutation effecting raffinose  
30 saccharides that was present in the original cross. Line 4E76 thus represents the LR33 mutation in a selected elite cultivar background. The line was grown in a total of nine environments over two years along with selected elite check lines for carbohydrate analysis. A more limited set of samples from three environments was analyzed for phytic acid content. The carbohydrate data is shown in Table 10 and the phytic acid  
35 data in Table 11.

Table 10

Mean, standard deviation (SD) and two standard deviations (2SD) of the mean for the stachyose, raffinose and sucrose content of LR33-containing line 4E76 and thirteen elite cultivar checks  
(All values are expressed as  $\mu\text{moles g}^{-1}$  seed weight.)

LINE	STACHYOSE			REFFINOSE			SUCROSE		
	MEAN	SD	2SD	MEAN	SD	2SD	MEAN	SD	2SD
4E76	2.4	0.5	1.0	6.9	0.7	1.4	249	42.9	85.8
A2514	44.8	2.7	5.4	13.0	2.6	5.2	138	9.6	19.2
A2396	47.8	1.1	2.2	9.6	0.9	1.8	158	9.6	19.2
A1923	50.2	2.8	5.6	8.6	0.5	1.0	148	9.4	18.8
A2506	51.0	3.7	7.4	16.4	4.0	8.0	161	28.2	56.4
A3322	54.2	7.3	14.6	14.0	2.7	5.4	147	27.1	54.2
A2923	56.4	2.2	4.4	13.6	1.7	3.4	125	9.2	18.4
A2234	57.2	9.1	18.2	16.6	3.4	6.8	145	21.2	42.4
A1923	57.5	4.5	9.0	18.5	3.4	6.8	156	29.8	59.6
A3313	61.0	9.7	19.4	14.4	3.9	7.8	172	37.9	75.8
A1662	61.6	6.2	12.4	13.8	3.4	6.8	127	21.4	42.8
A3935	62.8	9.2	18.4	11.6	2.1	4.2	184	38.7	77.4
A2833	63.8	8.6	17.2	15.4	2.4	4.8	150	38.7	77.4
A3510	68.8	9.4	18.8	14.2	1.9	3.8	160	33.0	66.0

Table 11

Mean, standard deviation and two standard deviations of the mean for the phytic acid content of the LR33-containing line 4E76 and the elite cultivar check A2872.  
(All values are expressed as  $\mu\text{moles g}^{-1}$  seed weight and represent data from three environments.)

LINE	PHYTIC ACID		
	MEAN	SD	2SD
4E76	12.3	2.4	4.8
A2872	20.9	0.9	1.8

In comparison to elite soybean cultivars that are typical of commercial soybeans, the LR33-derived line is eight to nine fold lower in the mass of total raffinose saccharide per gram of seed weight. The mass of phytic acid is also decreased by about 40%.

In comparison to elite soybean cultivars, the LR33-derived lines are lower in the mass of both raffinose and stachyose. While raffinose content is only slightly lower than values seen in elite lines, stachyose content is very greatly reduced. It is known that the detrimental effect of soybean raffinose saccharide content on energy



use efficiency when soybean meal is fed to monogastric animals is due to the poor digestibility of the  $\alpha$ -galactosidic bond. Accordingly, a decrease in combined raffinose plus stachyose content is an appropriate measure of increased digestibility. In fact, this measurement may even underestimate the effect of the instant phenotype since stachyose contains two moles of the  $\alpha$ -galactosidic bond per mole of sugar.

The absolute raffinose saccharide content of mature soybeans is known to vary in response to environmental factors. The effect of the mutation present in LR33-derived lines is of sufficient magnitude to render the combined raffinose plus stachyose content of such lines below 14.5  $\mu\text{mol/g}$  seed weight and should always remain far below that of wild type soybeans grown in the same environment.

The phytic acid content of mature soybean seeds is also known to vary in response to environmental factors, primarily due to variation of levels of available phosphate in soil. Once again, the mutation present in LR33-derived lines maintains total seed phytic acid content below 17  $\mu\text{mol/g}$  across all growth environments.

#### EXAMPLE 5

##### MOLECULAR IDENTIFICATION OF THE LR33 MUTATION

The evidence from analysis of metabolites in the LR33 derived lines presented in Example 2 strongly suggests that the LR33 mutation causes a decrease in the ability of the seed to produce *myo*-inositol.

In plants as well as other organisms, *myo*-inositol is produced solely by a pathway which begins with the conversion of glucose-6-phosphate to *myo*-inositol 1-phosphate in a reaction catalyzed by *myo*-inositol 1-phosphate synthase and ends with the hydrolysis of the 1-phosphate by a specific *myo*-inositol-phosphate phosphatase (see Figure 1; Loews, F.A. In: *Inositol Metabolism in Plants* (1990) Wiley-Liss, New York, pp 13-19)). Since the phytic acid has as its probable precursor *myo*-inositol 1-phosphate and since the level of phytic acid is decreased by the LR33 mutation, the gene and gene product for *myo*-inositol-phosphate synthase was characterized in both wild type and LR33 mutant plants.

##### cDNA cloning of the wild type soybean *myo*-inositol 1-phosphate synthase

A cDNA library was made as follows. Soybean embryos (ca. 50 mg fresh weight each) were removed from their pods and frozen in liquid nitrogen. The frozen embryos were ground to a fine powder in the presence of liquid nitrogen and then extracted by Polytron homogenization and fractionated to enrich for total RNA by the method of Chirgwin et al. ((1979) *Biochemistry* 18:5294-5299).

The nucleic acid fraction was enriched for poly A<sup>+</sup> RNA by passing total RNA through an oligo-dT cellulose column and eluting the poly A<sup>+</sup> RNA with salt as described by Goodman et al. ((1979) *Meth. Enzymol.* 68:75-90). cDNA was synthesized from the purified poly A<sup>+</sup> RNA using cDNA Synthesis System (Bethesda Research Laboratory, Gaithersburg, MD) and the manufacturer's instructions. The

resultant double-stranded DNA was methylated by Eco RI DNA methylase (Promega) prior to filling in its ends with T4 DNA polymerase (Bethesda Research Laboratory) and blunt-end ligation to phosphorylated Eco RI linkers using T4 DNA ligase (Pharmacia). The double-stranded DNA was digested with Eco RI enzyme, separated  
5 from excess linkers by passage through a gel filtration column (Sephadose CL-4B), and ligated to lambda ZAP vector (Stratagene) according to manufacturer's instructions. Ligated DNA was packaged into phage using the Gigapack™ packaging extract (Stratagene) according to manufacturer's instructions. The resultant cDNA library was amplified as per Stratagene's instructions and stored at -80°C.

10 Following the instructions in the Lambda ZAP Cloning Kit Manual (Stratagene), the cDNA phage library was used to infect *E. coli* XL1 cells and a total of approximately 300,000 plaque forming units were plated onto six 150 mm diameter petri plates.

Duplicate lifts of the plates were made onto nitrocellulose filters (Schleicher &  
15 Schuell, Keene, NH). The filters were prehybridized in 25 mL of hybridization buffer consisting of 6X SSPE, 5X Denhardt's solution, 0.5% SDS, 5% dextran sulfate and 0.1 mg/mL denatured salmon sperm DNA (Sigma Chemical Co.) at 60°C for 2 h.

The blocked filters were then hybridized to a radiolabelled probe made from a cDNA from *Arabidopsis thaliana* which had been identified as a *myo*-inositol-1-  
20 phosphate synthase by homology to yeast *myo*-inositol-1-phosphate synthase [Johnson, M.A. (1994) *Plant Physiol.* 105:1023-1024]. The *Arabidopsis* clone was obtained from the Arabidopsis Biological Resource Center, DNA Stock Center, 1060 Carmack Road, Columbus, OH 43210-1002, clone number 181C18T7113E9T7. The 1.2 kB cDNA insert was removed from the vector DNA by digestion with Sal I and  
25 Not I followed by agarose gel purification of the DNA fragment. The purified fragment was labeled with [<sup>32</sup>P]dCTP with a random primer labeling kit (Bethesda Research Laboratory). The filters were allowed to hybridize overnight under the same conditions as described for pre-hybridization. Excess radiolabel was washed from the filters in 0.6x SSC containing 0.1% SDS. Two washes of about 10 min each in  
30 0.2xSSC, 0.1% SDS at 60°C were then applied to remove non-specifically bound label and the filters were used to expose photographic film in an overnight exposure. Approximately 200 positive signals were observed. Six were purified by excision of the area around the signal, re-plating the phage and re-screening as above. Two clones were excised to phagmids and used to infect *E. coli* to obtain plasmid clones  
35 using the protocols described by the manufacturer (Stratagene). Of the two clones, one designated p5bmi-1-ps was sequenced using Applied Biological Instruments methodology and equipment. The nucleotide sequence of the cDNA insert in p5bmi-1-ps is shown in SEQ ID NO:1 and the deduced amino acid sequence encoded by that sequence in SEQ ID NO:2.

cDNA cloning of *myo*-inositol 1-phosphate synthase from  
immature seeds of LR33 soybeans

Seeds from the LR33 plants numbered 11, 16 and 20 and described in Example 3 (see Table 8) were harvested at about 50% through the seed filling period, removed from the pod and stored frozen at -80°C. For mRNA isolation, 2 g of frozen seed from the bulked seed population were ground in liquid nitrogen in a mortar and pestle.

The frozen powder was again ground in 10 mL of total RNA extraction buffer which consisted of 10 mM Tris-HCl, pH 9, 10 mM EDTA, 0.5% CTAB (cetyltrimethyl ammonium bromide), 0.8 M NaCl, 1% 2-mercaptoethanol, and 2% polyvinylpyrrolidone. Water for all reagents was treated with 0.05% diethylpyrocarbamate for 30 min then autoclaved. The tissue slurry was transferred to a 15 mL polypropylene tube and centrifuged at 5,500 x g for 15 min. The supernatant was passed through Miracloth (Calbiochem) into a second polypropylene tube and 0.3 volume of chloroform was added. The phases were separated by centrifugation at 5,500 x g for 10 min and the upper phase transferred to another polypropylene tube to which was added 1.5 volumes 10 mM Tris-HCl pH 9, 10 mM EDTA, 0.5% CTAB and 0.1% 2-mercaptoethanol. After a 30 min incubation at room temperature the nucleic acids were precipitated by centrifugation at 5,500 x g for 20 min.

The nucleic acids were re-dissolved in 0.4 mL of 1 M NaCl containing 0.1% 2-mercaptoethanol. After extraction with one volume of 1:1 phenol:chloroform, the nucleic acids were precipitated with two volumes of ethanol.

mRNA was purified from this nucleic acid fraction using the mRNA purification kit from Pharmacia. Approximately 12 µg of mRNA was obtained. Thirteen ng of the polyadenylated mRNA was used as template for amplification from oligo-dT using a GeneAmp® RNA-PCR kit (Perkin Elmer Cetus, part no. N808-0017). The reverse transcriptase reaction was run for 30 min at 42°C. For the PCR amplification, Vent™ DNA polymerase (New England Biolabs) was substituted for the DNA polymerase supplied by the kit manufacturer and an additional 2 µL of 100 mM magnesium sulfate was added to each 100 µL reaction. The 5' primer had the sequence shown in SEQ ID NO:3 and consists of bases 57 to 77 in SEQ ID NO:1 with the additional bases 5'-GGGAATTCCATATG-3' added to encode an Nde I site in the primer with eight additional 5' bases to enhance the restriction enzyme activity against the sequence.

The 3' primer had the sequence shown in SEQ ID NO:4 and consists of the reverse complement of bases 1566 to 1586 in SEQ ID NO:1 with the additional bases 5'-AAGGAAAAAGCGGCCGC-3' added to provide a Not I site in the primer and ten additional bases to enhance restriction digestion. The PCR reaction was run for 35 cycles at a 52°C annealing temperature and 1.5 min extension time. A product of

about 1550 base pairs was obtained and purified by passage through an Amicon 50 microfuge filter followed by extraction with an equal volume of 1:1 phenol:chloroform, extraction of the upper layer of the phenol:chloroform separation with one volume of chloroform and precipitation with ethanol. Five µg of the  
5 resulting, clean PCR product was digested overnight at 37°C with both Nde I and Not I. The restriction enzyme digest was de-proteinized by the above described phenol:chloroform extraction procedure and ligated into 2 µg of pET24aT7 expression vector (Novogen) that had also been digested with Nde I and Not I and treated with calf intestine alkaline phosphatase to hydrolyzed the terminal phosphates. The  
10 ligation mixture was used to transform electrocompetant DH 10B *E. coli* cells and transformants were selected by growth on plates containing 30 mg l<sup>-1</sup> kanamycin. Eighteen single colonies from the transformation plate were picked and placed in 100 µL of sterile water. Forty µL of the cell mix was used as the DNA template in a PCR reaction run with Taq™ polymerase (Perkin Elmer) using the primers and PCR  
15 conditions that were initially used to isolate the cDNA insert. Six colonies served as template to amplify a product of the correct size. The remaining 60 µL of the cell mix from these clones was grown in overnight culture and plasmid DNA preparations were made from each clone. The purified plasmid from each of the six clones was used to transform electrocompetant DE 3 *E. coli* cells.

20 Functional expression of the *myo*-inositol-1-phosphate synthase from wildtype and LR33 soybeans in *E. coli*

The wild type soybean *myo*-inositol 1-phosphate synthase was placed into the pET24aT7 expression vector by PCR amplification of p5bmi-1-ps using the primers in SEQ ID NO:3 and SEQ ID NO:4, and the PCR amplification and cloning protocol  
25 described above for cloning the LR33 *myo*-inositol 1-phosphate synthase from reverse transcribed mRNA. In this case, plasmid preparations from nine DH 10B clones that were shown to contain plasmid with the cDNA insert were pooled and used to transform electrocompetant DE 3 *E. coli* cells.

Kanamycin resistant colonies were selected and six were chosen for  
30 inoculation of overnight cultures. The overnight cultures grown at 30°C in LB media with 30 mg l<sup>-1</sup> kanamycin, were diluted two fold with fresh media, allowed to re-grow for 1 h, and induced by adding isopropyl-thiogalactoside to 1 mM final concentration.

Cells were harvested by centrifugation after 3 h and re-suspended in 50 µL of 50 mM Tris-HCl at pH 8.0 containing 0.1 mM DTT and 0.2 mM phenyl  
35 methylsulfonyl fluoride. A small amount of 1 mm glass beads were added and the mixture was sonicated three times for about five seconds each time with a microprobe sonicator. The mixture was centrifuged and the protein concentration of the supernatant was determined. One µg of protein from the soluble fraction of each

clonal culture was separated by SDS-PAGE. Cultures which produced an addition protein band of about 60 kilodaltons in mass were chosen for activity assay.

For the assay, [ $^{33}\text{P}$ ]glucose-6-phosphate was prepared by the hexokinase catalyzed phosphorylation of glucose using [ $^{33}\text{P}$ ]- $\gamma$ -ATP as the phosphate donor.

5 After 30 min at room temperature, the reaction mix was passed through a SEP-PAC C18 column (MilliPore Corp. Milford, MA) which had been washed first with 80% methanol and then water. The column pass through along with an additional 0.5 mL was collected and passed through a SEP-PAC SAX column which was then washed with 1 mL of 80% methanol. The [ $^{33}\text{P}$ ]-glucose-6-phosphate was eluted with 2 mL of  
10 0.02N HCl in 80% methanol. Forty  $\mu\text{L}$  of 1M Tris base was added and the methanol was removed under vacuum. The glucose-6-phosphate concentration of the remaining solution was determined by its conversion to 6-phosphoglucuronic acid by glucose-6-phosphate dehydrogenase. The NADPH produced in the reaction was quantitated by absorbance at 340 nm.

15 Ten  $\mu\text{L}$  of the cell extracts were incubated at 37°C for 30 min in 100  $\mu\text{L}$  reactions that were 2 mM in glucose-6-phosphate (57,334 dpm total  $^{33}\text{P}$ ), 0.1 mM in NAD, and 15 mM in ammonium acetate. The reactions were heated to 90°C to precipitate protein, centrifuged to clear. Forty-seven  $\mu\text{L}$  of the supernatant was applied to a Dionex<sup>TM</sup> PA-1 column run at 0.9 ml min<sup>-1</sup> with 0.1 N NaOH and 0.1 N  
20 sodium acetate. Standards of *myo*-inositol 1-phosphate eluted from 8 through 10 min after injection and 1 min fractions of the separated reaction mixes were taken through that time range for scintillation counting. The radioactivity in the peak fractions was summed to obtain the conversion of glucose-6-phosphate to *myo*-inositol-1-phosphate and the results for one control *E. coli* DE 3 culture containing an empty pET24aT7  
25 vector and two clones containing the soybean *myo*-inositol 1-phosphate synthase cDNA are shown in Table 12.

Table 12

The specific activity ( $\mu\text{moles myo-inositol 1-phosphate produce min}^{-1} \text{ mg protein}^{-1}$ ) for three *E. coli* cell culture extracts  
(Soy Clones 1 and 2 contain the soybean *myo*-inositol 1-phosphate synthase while the control contains an empty plasmid.)

LINE	SPECIFIC ACTIVITY
CONTROL	0.024 $\mu\text{mol min}^{-1} \text{ mg}^{-1}$
SOY CLONE 1	0.524 $\mu\text{mol min}^{-1} \text{ mg}^{-1}$
SOY CLONE 2	0.892 $\mu\text{mol min}^{-1} \text{ mg}^{-1}$

30 In the assay as run, both cDNA-containing clones used essentially all of the available substrate and were therefore more than 35 fold more active than the control

line. The soybean *myo*-inositol 1-phosphate synthase gene is therefore capable of producing a functional enzyme in *E. coli*.

The wild type soybean *myo*-inositol 1-phosphate synthase clone number 1 (Soy Clone 1) and three of the LR33 *myo*-inositol 1-phosphate synthase clones were grown for protein expression as described above. Five  $\mu\text{g}$  of protein from the soluble cell extract from each clone was separated by SDS-PAGE. LR33 clone number 10 (LR33-10) produced a soluble, 60 kilodalton protein in essentially the same abundance as did the wild type clone number 1. Four  $\mu\text{g}$  of protein from each of the two extracts was assayed for *myo*-inositol 1-phosphate synthase activity by the method described above using a 100  $\mu\text{L}$  reaction that was 3  $\mu\text{M}$  in NAD and 90  $\mu\text{M}$  in glucose-6-phosphate. The specific activity of the wild type *myo*-inositol 1-phosphate synthase was 1.5  $\text{nmol min}^{-1} \text{mg protein}^{-1}$  under these conditions while the specific activity of the LR33-derived *myo*-inositol 1-phosphate synthase was 0.16  $\text{nmol min}^{-1} \text{mg protein}^{-1}$ .

15 The nucleotide sequence of the cDNA encoding the LR33  
*myo*-inositol 1-phosphate synthase

The nucleotide sequence of the cDNA insert in clone LR33-10 (containing the nucleic acid fragment encoding the LR33 *myo*-inositol 1-phosphate synthase) was determined using the DH 10 B *E. coli* strain of that clone as the plasmid source and DNA sequencing as described for the wild type clone. The nucleotide sequence is shown in SEQ ID NO:5 and the deduced amino acid sequence obtained from the open reading frame of that clone in SEQ ID NO:6. SEQ ID NO:5 differs by single base pair change of G to T in the coding strand at base number 1241 from SEQ ID NO:1. That change results in a change of amino acid number 396 from lysine in the wild type sequence (SEQ ID NO:2) to asparagine in the LR33 amino acid sequence (SEQ ID NO:6).

To confirm that the base change resulted from a change in the LR33 genome rather than a PCR generated error which might have occurred during the cloning of the LR33 *myo*-inositol 1-phosphate synthase, two sets of PCR primers were prepared. The wild type primer (SEQ ID NO:7) and the LR33 primer (SEQ ID NO:8) were used to amplify genomic DNA prepared from dry seeds of soybean cultivar A2872 and from soybean line LR33 clone number 16 (see Table 8). The PCR primer described in SEQ ID NO:4 was used as the common antisense strand primer. At annealing temperatures of 62°C or 64°C and 35 cycles of annealing and extension, only the wild type primer produced a PCR product when A2872 DNA was used as a template and only the primer corresponding to the LR33 sequence produced a product when LR33 DNA was used as template.

To further check the specificity of the primers for detecting the mutation, DNA was prepared from six single plants grown from seed of the segregating LR33 clone

number 7 line described in Example 3 (see Table 8). Out of six plants tested from the segregating population, two gave DNA that acted as template for both primers, three gave DNA that acted as a primer only with the wild type primer, and one gave DNA that produced a product only with the LR33 primer. These are the results expected from a population that contains heterozygotes containing one wild type and one mutant copy of the gene.

From the metabolite data and sequence data, we conclude that the observed seed phenotype of very low total raffinose saccharide sugars, very high sucrose and low phytic acid are all due to the single base change mutation described by the comparison of the wild type and LR33 sequences shown in SEQ ID NO:1 and SEQ ID NO:5.

#### EXAMPLE 6

##### TRANSFORMATION OF SOYBEANS TO ACHIEVE GENE SILENCING OF *MYO*-INOSITOL-1-PHOSPHATE-SYNTHASE AND THE ASSOCIATED SEED PHENOTYPE

##### Construction of vectors for transformation of *Glycine max* for reduced expression of *myo*-inositol 1-phosphate synthase in developing soybean seeds

Plasmids containing the antisense or sense oriented soybean *myo*-inositol 1-phosphate synthase cDNA sequence under control of the soybean Kunitz Trypsin Inhibitor 3 (KTI3) promoter [Jofuku and Goldberg, (1989) *Plant Cell* 1:1079-1093], the *Phaseolus vulgaris* 7S seed storage protein (phaseolin) promoter [Sengupta-Gopalan et al., (1985) *Proc. Natl. Acad. Sci. USA* 82:3320-3324; Hoffman et al., (1988) *Plant Mol. Biol.* 11:717-729] and soybean  $\beta$ -conglycinin promoter [Beachy et al., (1985) *EMBO J.* 4:3047-3053], are constructed. The construction of vectors expressing the soybean *myo*-inositol-1-phosphate synthase cDNA under the control of these promoters is facilitated by the use of the following plasmids: pML70, pCW108 and pCW109A.

The pML70 vector contains the KTI3 promoter and the KTI3 3' untranslated region and was derived from the commercially available vector pTZ18R (Pharmacia) via the intermediate plasmids pML51, pML55, pML64 and pML65. A 2.4 kb Bst BI/Eco RI fragment of the complete soybean KTI3 gene [Jofuku and Goldberg *supra*], which contains all 2039 nucleotides of the 5' untranslated region and 390 bases of the coding sequence of the KTI3 gene ending at the Eco RI site corresponding to bases 755 to 761 of the sequence described in Jofuku et al., (1989) *Plant Cell* 1:427-435, was ligated into the Acc I/Eco RI sites of pTZ18R to create the plasmid pML51. The plasmid pML51 was cut with Nco I, filled in using Klenow, and religated, to destroy an Nco I site in the middle of the 5' untranslated region of the KTI3 insert, resulting in the plasmid pML55. The plasmid pML55 was partially digested with Xmn I/Eco RI to release a 0.42 kb fragment, corresponding to bases 732 to 755 of the above cited

sequence, which was discarded. A synthetic Xmn I/Eco RI linker containing an Nco I site, was constructed by making a dimer of complementary synthetic oligonucleotides consisting of the coding sequence for an Xmn I site (5'-TCTTCC-3') and an Nco I site (5'-CCATGGG-3') followed directly by part of an Eco RI site (5'-GAAGG-3').

- 5 The Xmn I and Nco I/Eco RI sites were linked by a short intervening sequence (5'-ATAGCCCCCAA-3'). This synthetic linker was ligated into the Xmn I/Eco RI sites of the 4.94 kb fragment to create the plasmid pML64. The 3' untranslated region of the KTi3 gene was amplified from the sequence described in Jofuku et al., [*supra*] by standard PCR protocols (Perkin Elmer Cetus, GeneAmp PCR kit) using the
- 10 primers ML51 and ML52. Primer ML51 contained the 20 nucleotides corresponding to bases 1072 to 1091 of the above cited sequence with the addition of nucleotides corresponding to Eco RV (5'-GATATC-3'), Nco I (5'-CCATGG-3'), Xba I (5'-TCTAGA-3'), Sma I (5'-CCCGGG-3') and Kpn I (5'-GGTACC-3') sites at the 5' end of the primer. Primer ML52 contained the exact complement of the nucleotides
- 15 corresponding to bases 1242 to 1259 of the above cited sequence with the addition of nucleotides corresponding to Sma I (5'-CCCGGG-3'), Eco RI (5'-GAATTC-3'), Bam HI (5'-GGATCC-3') and Sal I (5'-GTCGAC-3') sites at the 5' end of the primer. The PCR-amplified 3' end of the KTi3 gene was ligated into the Nco I/Eco RI sites of pML64 to create the plasmid pML65. A synthetic multiple cloning site linker was
- 20 constructed by making a dimer of complementary synthetic oligonucleotides consisting of the coding sequence for Pst I (5'-CTGCA-3'), Sal I (5'-GTCGAC-3'), Bam HI (5'-GGATCC-3') and Pst I (5'-CTGCA-3') sites. The linker was ligated into the Pst I site (directly 5' to the KTi3 promoter region) of pML65 to create the plasmid pML70.
- 25 The pCW108 vector contains the bean phaseolin promoter and 3' untranslated region and was derived from the commercially available pUC18 plasmid (Gibco-BRL) via plasmids AS3 and pCW104. Plasmid AS3 contains 495 base pairs of the bean (*Phaseolus vulgaris*) phaseolin (7S seed storage protein) promoter starting with 5'-TGGTCTTTTGGT-3' followed by the entire 1175 base pairs of the 3'
- 30 untranslated region of the same gene [see sequence descriptions in Doyle et al., (1986) *J. Biol. Chem.* 261:9228-9238 and Slightom et al., (1983) *Proc. Natl. Acad. Sci. USA* 80:1897-1901; further sequence description may be found in World Patent Publication WO911/3993] cloned into the Hind III site of pUC18. The additional cloning sites of the pUC18 multiple cloning region (Eco RI, Sph I, Pst I and Sal I) were removed by
- 35 digesting with Eco RI and Sal I, filling in the ends with Klenow and religating to yield the plasmid pCW104. A new multiple cloning site was created between the 495 bp of the 5' phaseolin and the 1175 bp of the 3' phaseolin by inserting a dimer of complementary synthetic oligonucleotides consisting of the coding sequence for a Nco I site (5'-CCATGG-3') followed by three filler bases (5'-TAG-3'), the coding



sequence for a Sma I site (5'-CCCGGG-3'), the last three bases of a Kpn I site (5'-TAC-3'), a cytosine and the coding sequence for an Xba I site (5'-TCTAGA-3') to create the plasmid pCW108. This plasmid contains unique Nco I, Sma I, Kpn I and Xba I sites directly behind the phaseolin promoter.

5       The vector pCW109A contains the soybean  $\beta$ -conglycinin promoter sequence and the phaseolin 3' untranslated region and is a modified version of vector pCW109 which was derived from the commercially available plasmid pUC18 (Gibco-BRL). The vector pCW109 was made by inserting into the Hind III site of the cloning vector pUC18 a 555 bp 5' non-coding region (containing the promoter region) of the  
10        $\beta$ -conglycinin gene followed by the multiple cloning sequence containing the restriction endonuclease sites for Nco I, Sma I, Kpn I and Xba I, as described for pCW108 above, then 1174 bp of the common bean phaseolin 3' untranslated region into the Hind III site (described above).

15       The  $\beta$ -conglycinin promoter region used is an allele of the published  $\beta$ -conglycinin gene [Doyle et al., (1986) *J. Biol. Chem.* 261:9228-9238] due to differences at 27 nucleotide positions. Further sequence description of this gene may be found in World Patent Publication W091/13993.

20       These three nucleic acid constructions constitute seed specific expression vectors with expression over a wide developmental period including the period of *myo*-inositol synthesis for subsequent conversion to phytic acid. Insertion of the sequences described in SEQ ID NO:1 and SEQ ID NO:5 into these vectors is facilitated by the PCR methods described in Example 5 above. PCR primers which are complementary to chosen regions of SEQ ID NO:1 or SEQ ID NO:5 may be synthesized with additional bases that constitute the recognition sequences of  
25       restriction endonucleases chosen from among those that also cut in the multiple cloning sequences following the promoter sequences of pML70, pCW108 and pCW109. Placement of the restriction sites may be chosen so as to direct the orientation of the nucleotide fragment from the soybean *myo*-inositol 1-phosphate synthase into the sense orientation to achieve either over expression or co-suppression  
30       or into the antisense orientation to achieve under expression.

Transformation Of Somatic Soybean Embryo Cultures  
and Regeneration Of Soybean Plants

35       The following stock solutions and media are used to support the growth of soybean tissues *in vitro*:

Stock Solutions:

MS Sulfate (100X Stock)		MS Halides (100X stock)	
(g/L)		(g/L)	
MgSO <sub>4</sub> 7H <sub>2</sub> O	37.0	CaCl <sub>2</sub> 2H <sub>2</sub> O	44.0
MnSO <sub>4</sub> H <sub>2</sub> O	1.69	KI	0.083
ZnSO <sub>4</sub> 7H <sub>2</sub> O	0.86	CoCl <sub>2</sub> 6H <sub>2</sub> O	0.00125
CuSO <sub>4</sub> 5H <sub>2</sub> O	0.0025	KH <sub>2</sub> PO <sub>4</sub>	17.0
		H <sub>3</sub> BO <sub>3</sub>	0.63
		Na <sub>2</sub> MoO <sub>4</sub> 2H <sub>2</sub> O	0.025

B5 Vitamin		MS FeEDTA (100X stock)	
(g/L)		(g/L)	
<i>myo</i> -inositol	10.0	Na <sub>2</sub> EDTA	3.72
nicotinic acid	0.10	FeSO <sub>4</sub> 7H <sub>2</sub> O	2.784
pyridoxine HCl	0.10		

Media:

Component (per liter)	SB55	SBP6	SB103	SB71-1
MS Sulfate Stock	10 mL	10 mL	10 mL	-
MS Halides Stock	10 mL	10 mL	10 mL	-
MS FeEDTA Stock	10 mL	10 mL	10 mL	-
B5 Vitamin Stock	1 mL	1 mL	-	1 mL
2,4-D Stock (10 mg/ml)	1 mL	0.5 mL	-	-
Sucrose	60 g	60 g	-	3%
Maltose	-	-	6%	-
Asparagine	0.667 g	0.667 g	-	-
NH <sub>4</sub> NO	0.8 g	0.8 g	-	-
KNO <sub>3</sub>	3.033g	3.033g	-	-
MgCl <sub>2</sub>	-	-	750 mg	750 mg
Gelrite	-	-	0.2%	0.2%
pH	5.7	5.7	5.7	5.7

5

Soybean embryogenic suspension cultures are maintained in 35 mL liquid media (SB55 or SBP6) on a rotary shaker, 150 rpm, at 28°C with mixed florescent and incandescent lights on a 16:8 h day/night schedule. Cultures are subcultured every four weeks by inoculating approximately 35 mg of tissue into 35 mL of liquid medium.

10

Soybean embryogenic suspension cultures may be transformed with seed specific expression vectors containing either the sense or antisense oriented

*myo*-inositol 1-phosphate synthase by the method of particle gun bombardment (see Kline et al. (1987) *Nature (London)* 327:70). A DuPont Biolistic™ PDS1000/HE instrument (helium retrofit) is used for these transformations.

To 50 mL of a 60 mg/mL 1 µm gold particle suspension are added (in order):  
5 5 µL DNA (1 µg/µL), 20 µL spermidine (0.1 M), and 50 µl CaCl<sub>2</sub> (2.5 M). The particle preparation is agitated for 3 min, spun in a microfuge for 10 sec and the supernatant removed. The DNA-coated particles are then washed once in 400 µL 70% ethanol and resuspended in 40 µL of anhydrous ethanol. The DNA/particle suspension is sonicated three times for 1 sec each. Five µL of the DNA-coated gold  
10 particles are then loaded on each macro carrier disk.

Approximately 300-400 mg of a four week old suspension culture is placed in an empty 60 x 15 mm petri dish and the residual liquid removed from the tissue with a pipette. For each transformation experiment, approximately 5-10 plates of tissue were normally bombarded. Membrane rupture pressure is set at 1000 psi and the chamber  
15 is evacuated to a vacuum of 28 inches of mercury. The tissue is placed approximately 3.5 inches away from the retaining screen and bombarded three times. Following bombardment, the tissue is placed back into liquid and cultured as described above.

Eleven days post bombardment, the liquid media is exchanged with fresh SB55 containing 50 mg/mL hygromycin. Thereafter, the selective media is refreshed  
20 weekly. Seven weeks post bombardment, green, transformed tissue may be observed growing from untransformed, necrotic embryogenic clusters. Isolated green tissue is removed and inoculated into individual flasks to generate new, clonally propagated, transformed embryogenic suspension cultures. Thus each new line is treated as independent transformation event. These suspensions can then be maintained as  
25 suspensions of embryos clustered in an immature developmental stage through subculture or regenerated into whole plants by maturation and germination of individual somatic embryos.

Transformed embryogenic clusters are removed from liquid culture and placed on a solid agar media (SB103) containing no hormones or antibiotics. Embryos are  
30 cultured for eight weeks at 26°C with mixed florescent and incandescent lights on a 16:8 h day/night schedule. During this period, individual embryos can be removed from the clusters and analyzed at various stages of embryo development. After eight weeks somatic embryos become suitable for germination. For germination, eight week old embryos are removed from the maturation medium and dried in empty petri  
35 dishes for 1 to 5 days. The dried embryos are then planted in SB71-1 medium where they are allowed to germinate under the same lighting and germination conditions described above. Germinated embryos can then be transferred to sterile soil and grown to maturity for seed collection.

While in the globular embryo state in liquid culture as described above, somatic soybean embryos contain very low amounts of triacylglycerol or storage proteins typical of maturing, zygotic soybean embryos. At this developmental stage, the ratio of total triacylglyceride to total polar lipid (phospholipids and glycolipid) is about 1:4, as is typical of zygotic soybean embryos at the developmental stage from which the somatic embryo culture was initiated. At the globular stage as well, the mRNAs for the prominent seed proteins ( $\alpha'$ -subunit of  $\beta$ -conglycinin, Kunitz Trypsin Inhibitor 3 and Soybean Seed Lectin) are essentially absent. Upon transfer to hormone-free media to allow differentiation to the maturing somatic embryo state as described above, triacylglycerol becomes the most abundant lipid class. As well, mRNAs for  $\alpha'$ -subunit of  $\beta$ -conglycinin, Kunitz Trypsin Inhibitor 3 and Soybean Seed Lectin become very abundant messages in the total mRNA population. In these respects the somatic soybean embryo system behaves very similarly to maturing zygotic soybean embryos *in vivo*. During the early maturation period the main soluble carbohydrates present in the somatic embryos are sucrose, glucose and maltose (the supplied sugar during the maturation phase). As the somatic embryos mature and begin to yellow, both raffinose and stachyose are formed. In this respect as well the phenotype of the somatic embryos is very similar to zygotic embryos as they go through the late stages of seed development. Thus selection for embryos that are transformed with seed specific expression vectors which direct the expression of the nucleotide sequences described in SEQ ID NO:1 or SEQ ID NO:5 I in either the sense or antisense orientation and that produce reduced amount of raffinose and stachyose should lead to the regeneration of mature, fertile soybean plants which bear seeds with that same phenotype.

25

SEQUENCE LISTING

## (1) GENERAL INFORMATION:

- (i) APPLICANT: HITZ, WILLIAM D.  
SEBASTIAN, SCOTT ANTHONY
- (ii) TITLE OF INVENTION: SOYBEAN PLANTS PRODUCING SEEDS WITH  
REDUCED LEVELS OF RAFFINOSE  
SACCHARIDES AND PHYTIC ACID
- (iii) NUMBER OF SEQUENCES: 8
- (iv) CORRESPONDENCE ADDRESS:
  - (A) ADDRESSEE: E. I. DU PONT DE NEMOURS AND COMPANY
  - (B) STREET: 1007 MARKET STREET
  - (C) CITY: WILMINGTON
  - (D) STATE: DELAWARE
  - (E) COUNTRY: UNITED STATES OF AMERICA
  - (F) ZIP: 19898
- (v) COMPUTER READABLE FORM:
  - (A) MEDIUM TYPE: DISKETTE, 3.50 INCH
  - (B) COMPUTER: IBM PC COMPATIBLE
  - (C) OPERATING SYSTEM: MICROSOFT WINDOWS 95
  - (D) SOFTWARE: MICROSOFT WORD FOR WINDOWS 95 (7.0)
- (vi) CURRENT APPLICATION DATA:
  - (A) APPLICATION NUMBER:
  - (B) FILING DATE:
  - (C) CLASSIFICATION:
- (viii) ATTORNEY/AGENT INFORMATION:
  - (A) NAME: MAJARIAN, WILLIAM R.
  - (B) REGISTRATION NUMBER: P41,173
  - (C) REFERENCE/DOCKET NUMBER: BB-1077
- (ix) TELECOMMUNICATION INFORMATION:
  - (A) TELEPHONE: (302)992-4926
  - (B) TELEFAX: (302)773-0164

## (2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1782 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: double
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA
- (iii) HYPOTHETICAL: NO
- (iv) ANTI-SENSE: NO
- (vi) ORIGINAL SOURCE:
  - (A) ORGANISM: Soybean line LR13
- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: p5bmi-1-ps
- (ix) FEATURE:
  - (A) NAME/KEY: CDS
  - (B) LOCATION: 54..1586
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

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TCGAGAATTT TAAGGTTGAG TGTCCTAATG TGAAGTACAC CGAGACTGAG ATTCAGTCCG    120
TGTACAAC TA CGAAACCACC GAACTTGTTT ACGAGAACAG GAATGGCACC TATCAGTGGA    180
TTGTCAAACC CAAATCTGTC AAATACGAAT TAAAACCAA CATCCATGTT CCTAAATTAG    240
GGGTAATGCT TGTGGGTTGG GGTGGAAACA ACGGCTCAAC CCTCACC GGTGTTATTG    300
CTAACCGAGA GGGCATTTC TGGGCTACAA AGGACAAGAT TCAACAAGCC AATTACTTTG    360
GCTCCCTCAC CCAAGCCTCA GCTATCCGAG TTGGGTCCTT CCAGGGAGAG GAAATCTATG    420
CCCCATTCAA GAGCCTGCTT CCAATGGTTA ACCCTGACGA CATTGTGTTT GGGGGATGGG    480
ATATCAGCAA CATGAACCTG GCTGATGCCA TGGCCAGGGC AAAGGTGTTT GACATCGATT    540
TGCAGAAGCA GTTGAGGCCT TACATGGAAT CCATGCTTCC ACTCCCCGGA ATCTATGACC    600
CGGATTTT CAT TGCTGCCAAC CAAGAGGAGC GTGCCAACA CGTCATCAAG GGCACAAAGC    660
AAGAGCAAGT TCAACAAATC ATCAAAGACA TCAAGGCGTT TAAGGAAGCC ACCAAAGTGG    720
ACAAGGTGGT TGTACTGTGG ACTGCCAACA CAGAGAGGTA CAGTAATTTG GTTGTGGGCC    780
TTAATGACAC CATGGAGAAT CTCTTGCTG CTGTGGACAG AAATGAGGCT GAGATTTCTC    840
CTTCCACCTT GTATGCCATT GCTTGTGTTA TGGAAAATGT TCCTTTCATT AATGGAAGCC    900
CTCAGAACAC TTTTGTACCA GGGCTGATTG ATCTTGCCAT CGCGAGGAAC ACTTTGATTG    960

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GTGGAGATGA CTTCAAGAGT GGTCAAGACCA AAATGAAATC TGTGTTGGTT GATTTCCTTG 1020  
 TGGGGGCTGG TATCAAGCCA ACATCTATAG TCAGTTACAA CCATCTGGGA AACAAATGATG 1080  
 GTATGAATCT TTCGGCTCCA CAAACTTTCC GTTCCAAGGA AATCTCCAAG AGCAACGTTG 1140  
 TTGATGATAT GGTCAACAGC AATGCCATCC TCTATGAGCC TGGTGAACAT CCAGACCATG 1200  
 TTGTTGTTAT TAAGTATGTG CCTTACGTAG GGGACAGCAA GAGAGCCATG GATGAGTACA 1260  
 CTTCAAGAGAT ATTCATGGGT GGAAAGAGCA CCATTGTTTT GCACAACACA TGCGAGGATT 1320  
 CCTCTTAGC TGCTCCTATT ATCTTGACT TGGTCCTTCT TGCTGAGCTC AGCACTAGAA 1380  
 TCGAGTTTAA AGCTGAAAT GAGGAAAAAT TCCACTCATT CCACCCAGTT GCTACCATCC 1440  
 TCAGCTACCT CACCAAGGCT CCTCTGGTTC CACCGGTAC ACCAGTGGTG AATGCATTGT 1500  
 CAAAGCAGCG TGCAATGCTG GAAACATAA TGAGGGCTTG TGTGGATTG GCGCCAGAGA 1560  
 ATAACATGAT TCTCGAGTAC AAGTGAAGCA TGGGACCGAA GAATAATATA GTTGGGGTAG 1620  
 CCTAGCTGAA TGTTTTATGT TAATAATATG TTTGCTTATA ATTTTGCAAG TGTAATTGAA 1680  
 TGCATCAGCT TCATTAATGC TTTAGAGCGG GGCATATTCT GTTTACTAGG AACATGAATG 1740  
 AATGTAGTAT AATTTTGTGT AAAAAAAAAA AAAAAAAAAA AA 1782

## (2) INFORMATION FOR SEQ ID NO:2:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 510 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

Met	Phe	Ile	Glu	Asn	Phe	Lys	Val	Glu	Cys	Pro	Asn	Val	Lys	Tyr	Thr
1				5				10					15		
Glu	Thr	Glu	Ile	Gln	Ser	Val	Tyr	Asn	Tyr	Glu	Thr	Thr	Glu	Leu	Val
			20					25					30		
His	Glu	Asn	Arg	Asn	Gly	Thr	Tyr	Gln	Trp	Ile	Val	Lys	Pro	Lys	Ser
			35				40					45			
Val	Lys	Tyr	Glu	Phe	Lys	Thr	Asn	Ile	His	Val	Pro	Lys	Leu	Gly	Val
			50			55					60				
Met	Leu	Val	Gly	Trp	Gly	Gly	Asn	Asn	Gly	Ser	Thr	Leu	Thr	Gly	Gly
65				70					75					80	
Val	Ile	Ala	Asn	Arg	Glu	Gly	Ile	Ser	Trp	Ala	Thr	Lys	Asp	Lys	Ile
			85					90						95	

Gln Gln Ala Asn Tyr Phe Gly Ser Leu Thr Gln Ala Ser Ala Ile Arg  
 100 105 110  
 Val Gly Ser Phe Gln Gly Glu Glu Ile Tyr Ala Pro Phe Lys Ser Leu  
 115 120 125  
 Leu Pro Met Val Asn Pro Asp Asp Ile Val Phe Gly Gly Trp Asp Ile  
 130 135 140  
 Ser Asn Met Asn Leu Ala Asp Ala Met Ala Arg Ala Lys Val Phe Asp  
 145 150 155 160  
 Ile Asp Leu Gln Lys Gln Leu Arg Pro Tyr Met Glu Ser Met Leu Pro  
 165 170 175  
 Leu Pro Gly Ile Tyr Asp Pro Asp Phe Ile Ala Ala Asn Gln Glu Glu  
 180 185 190  
 Arg Ala Asn Asn Val Ile Lys Gly Thr Lys Gln Glu Gln Val Gln Gln  
 195 200 205  
 Ile Ile Lys Asp Ile Lys Ala Phe Lys Glu Ala Thr Lys Val Asp Lys  
 210 215 220  
 Val Val Val Leu Trp Thr Ala Asn Thr Glu Arg Tyr Ser Asn Leu Val  
 225 230 235 240  
 Val Gly Leu Asn Asp Thr Met Glu Asn Leu Leu Ala Ala Val Asp Arg  
 245 250 255  
 Asn Glu Ala Glu Ile Ser Pro Ser Thr Leu Tyr Ala Ile Ala Cys Val  
 260 265 270  
 Met Glu Asn Val Pro Phe Ile Asn Gly Ser Pro Gln Asn Thr Phe Val  
 275 280 285  
 Pro Gly Leu Ile Asp Leu Ala Ile Ala Arg Asn Thr Leu Ile Gly Gly  
 290 295 300  
 Asp Asp Phe Lys Ser Gly Gln Thr Lys Met Lys Ser Val Leu Val Asp  
 305 310 315 320  
 Phe Leu Val Gly Ala Gly Ile Lys Pro Thr Ser Ile Val Ser Tyr Asn  
 325 330 335  
 His Leu Gly Asn Asn Asp Gly Met Asn Leu Ser Ala Pro Gln Thr Phe  
 340 345 350  
 Arg Ser Lys Glu Ile Ser Lys Ser Asn Val Val Asp Asp Met Val Asn  
 355 360 365  
 Ser Asn Ala Ile Leu Tyr Glu Pro Gly Glu His Pro Asp His Val Val  
 370 375 380  
 Val Ile Lys Tyr Val Pro Tyr Val Gly Asp Ser Lys Arg Ala Met Asp  
 385 390 395 400



Glu Tyr Thr Ser Glu Ile Phe Met Gly Gly Lys Ser Thr Ile Val Leu  
                     405                    410                    415  
 His Asn Thr Cys Glu Asp Ser Leu Leu Ala Ala Pro Ile Ile Leu Asp  
                     420                    425                    430  
 Leu Val Leu Leu Ala Glu Leu Ser Thr Arg Ile Glu Phe Lys Ala Glu  
                     435                    440                    445  
 Asn Glu Gly Lys Phe His Ser Phe His Pro Val Ala Thr Ile Leu Ser  
                     450                    455                    460  
 Tyr Leu Thr Lys Ala Pro Leu Val Pro Pro Gly Thr Pro Val Val Asn  
                     465                    470                    475                    480  
 Ala Leu Ser Lys Gln Arg Ala Met Leu Glu Asn Ile Met Arg Ala Cys  
                     485                    490                    495  
 Val Gly Leu Ala Pro Glu Asn Asn Met Ile Leu Glu Tyr Lys  
                     500                    505                    510

## (2) INFORMATION FOR SEQ ID NO:3:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 35 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (oligonucleotide)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

GGGAATTCCA TATGTTTCATC GAGAATTTTA AGGTT

35

## (2) INFORMATION FOR SEQ ID NO:4:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 39 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (oligonucleotide)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

AAGGAAAAAA GCGGCCGCTC ACTTGTA CTG GAGAATCAT

39

## (2) INFORMATION FOR SEQ ID NO:5:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 1775 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: double
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA

- (iii) HYPOTHETICAL: NO
- (iv) ANTI-SENSE: NO
- (vi) ORIGINAL SOURCE:  
(A) ORGANISM: Soybean line LR33
- (vii) IMMEDIATE SOURCE:  
(B) CLONE: LR33-10
- (ix) FEATURE:  
(A) NAME/KEY: CDS  
(B) LOCATION: 1..1533
- (ix) FEATURE:  
(A) NAME/KEY: POINT MUTATION  
(B) LOCATION: 1188
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

```

ATGTTTCATCG AGAATTTTAA GGTGAGTGT CCTAATGTGA AGTACACCGA GACTGAGATT    60
CAGTCCGTGT ACAACTACGA AACCACCGAA CTTGTTCACG AGAACAGGAA TGGCACCTAT    120
CAGTGGATTG TCAAACCCAA ATCTGTCAAA TACGAATTTA AAACCAACAT CCATGTTCCCT    180
AAATTAGGGG TAATGCTTGT GGGTTGGGGT GGAAACAACG GCTCAACCCT CACCGGTGGT    240
GTTATTGCTA ACCGAGAGGG CATTTCATGG GCTACAAAGG ACAAGATTCA ACAAGCCAAT    300
TACTTTGGCT CCCTCACCCA AGCCTCAGCT ATCCGAGTTG GGTCTTCCA GGGAGAGGAA    360
ATCTATGCCC CATTCAAGAG CCTGCTTCCA ATGGTTAACC CTGACGACAT TGTGTTTGGG    420
GGATGGGATA TCAGCAACAT GAACCTGGCT GATGCCATGG CCAGGGCAAA GGTGTTTGAC    480
ATCGATTTCG AGAAGCAGTT GAGGCCTTAC ATGGAATCCA TGCTTCCACT CCCCGBAATC    540
TATGACCCGG ATTCATTGCT TGCCAACCAA GAGGAGCGTG CCAACAACGT CATCAAGGGC    600
ACAAAGCAAG AGCAAGTTCA ACAAATCATC AAAGACATCA AGGCGTTTAA GGAAGCCACC    660
AAAGTGGACA AGGTGGTTGT ACTGTGGACT GCCAACACAG AGAGGTACAG TAATTTGGTT    720
GTGGGCCTTA ATGACACCAT GGAGAATCTC TTGGCTGCTG TGGACAGAAA TGAGGCTGAG    780
ATTTCTCCTT CCACCTTGTA TGCCATTGCT TGTGTTATGG AAAATGTTCC TTTCATTAAT    840
GGAAGCCCTC AGAACACTTT TGTACCAGGG CTGATTGATC TTGCCATCGC GAGGAACACT    900
TGATTGGTG GAGATGACTT CAAGAGTGGT CAGACCAAAA TGAAATCTGT GTTGGTTGAT    960
TTCCTTGTTG GGGCTGGTAT CAAGCCAACA TCTATAGTCA GTTACAACCA TCTGGGAAAC   1020
AATGATGGTA TGAATCTTTC GGCTCCACAA ACTTTCCGTT CCAAGGAAAT CTCCAAGAGC   1080
AACGTTGTTG ATGATATGGT CAACAGCAAT GCCATCCTCT ATGAGCCTGG TGAACATCCA   1140

```

GACCATGTTG TTGTTATTAA GTATGTGCCT TACGTAGGGG ACAGCAATAG AGCCATGGAT 1200  
 GAGTACACTT CAGAGATATT CATGGGTGGA AAGAGCACCA TTGTTTTGCA CAACACATGC 1260  
 GAGGATTCCC TCTTAGCTGC TCCTATTATC TTGGACTTGG TCCTTCTTGC TGAGCTCAGC 1320  
 ACTAGAATCG AGTTTAAAGC TGAAAATGAG GGAAAATTCC ACTCATTCCA CCCAGTTGCT 1380  
 ACCATCCTCA GCTACCTCAC CAAGGCTCCT CTGGTTCCAC CGGGTACACC AGTGGTGAAT 1440  
 GCATTGTCAA AGCAGCGTGC AATGCTGGAA AACATAATGA GGGCTTGTGT TGGATTGGCC 1500  
 CCAGAGAATA ACATGATTCT CGAGTACAAG TGA 1533

## (2) INFORMATION FOR SEQ ID NO:6:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 510 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

Met	Phe	Ile	Glu	Asn	Phe	Lys	Val	Glu	Cys	Pro	Asn	Val	Lys	Tyr	Thr	1	5	10	15
Glu	Thr	Glu	Ile	Gln	Ser	Val	Tyr	Asn	Tyr	Glu	Thr	Thr	Glu	Leu	Val	20	25	30	
His	Glu	Asn	Arg	Asn	Gly	Thr	Tyr	Gln	Trp	Ile	Val	Lys	Pro	Lys	Ser	35	40	45	
Val	Lys	Tyr	Glu	Phe	Lys	Thr	Asn	Ile	His	Val	Pro	Lys	Leu	Gly	Val	50	55	60	
Met	Leu	Val	Gly	Trp	Gly	Gly	Asn	Asn	Gly	Ser	Thr	Leu	Thr	Gly	Gly	65	70	75	80
Val	Ile	Ala	Asn	Arg	Glu	Gly	Ile	Ser	Trp	Ala	Thr	Lys	Asp	Lys	Ile	85	90	95	
Gln	Gln	Ala	Asn	Tyr	Phe	Gly	Ser	Leu	Thr	Gln	Ala	Ser	Ala	Ile	Arg	100	105	110	
Val	Gly	Ser	Phe	Gln	Gly	Glu	Glu	Ile	Tyr	Ala	Pro	Phe	Lys	Ser	Leu	115	120	125	
Leu	Pro	Met	Val	Asn	Pro	Asp	Asp	Ile	Val	Phe	Gly	Gly	Trp	Asp	Ile	130	135	140	
Ser	Asn	Met	Asn	Leu	Ala	Asp	Ala	Met	Ala	Arg	Ala	Lys	Val	Phe	Asp	145	150	155	160

Ile Asp Leu Gln Lys Gln Leu Arg Pro Tyr Met Glu Ser Met Leu Pro  
 165 170 175  
 Leu Pro Gly Ile Tyr Asp Pro Asp Phe Ile Ala Ala Asn Gln Glu Glu  
 180 185 190  
 Arg Ala Asn Asn Val Ile Lys Gly Thr Lys Gln Glu Gln Val Gln Gln  
 195 200 205  
 Ile Ile Lys Asp Ile Lys Ala Phe Lys Glu Ala Thr Lys Val Asp Lys  
 210 215 220  
 Val Val Val Leu Trp Thr Ala Asn Thr Glu Arg Tyr Ser Asn Leu Val  
 225 230 235 240  
 Val Gly Leu Asn Asp Thr Met Glu Asn Leu Leu Ala Ala Val Asp Arg  
 245 250 255  
 Asn Glu Ala Glu Ile Ser Pro Ser Thr Leu Tyr Ala Ile Ala Cys Val  
 260 265 270  
 Met Glu Asn Val Pro Phe Ile Asn Gly Ser Pro Gln Asn Thr Phe Val  
 275 280 285  
 Pro Gly Leu Ile Asp Leu Ala Ile Ala Arg Asn Thr Leu Ile Gly Gly  
 290 295 300  
 Asp Asp Phe Lys Ser Gly Gln Thr Lys Met Lys Ser Val Leu Val Asp  
 305 310 315 320  
 Phe Leu Val Gly Ala Gly Ile Lys Pro Thr Ser Ile Val Ser Tyr Asn  
 325 330 335  
 His Leu Gly Asn Asn Asp Gly Met Asn Leu Ser Ala Pro Gln Thr Phe  
 340 345 350  
 Arg Ser Lys Glu Ile Ser Lys Ser Asn Val Val Asp Asp Met Val Asn  
 355 360 365  
 Ser Asn Ala Ile Leu Tyr Glu Pro Gly Glu His Pro Asp His Val Val  
 370 375 380  
 Val Ile Lys Tyr Val Pro Tyr Val Gly Asp Ser Asn Arg Ala Met Asp  
 385 390 395 400  
 Glu Tyr Thr Ser Glu Ile Phe Met Gly Gly Lys Ser Thr Ile Val Leu  
 405 410 415  
 His Asn Thr Cys Glu Asp Ser Leu Leu Ala Ala Pro Ile Ile Leu Asp  
 420 425 430  
 Leu Val Leu Leu Ala Glu Leu Ser Thr Arg Ile Glu Phe Lys Ala Glu  
 435 440 445  
 Asn Glu Gly Lys Phe His Ser Phe His Pro Val Ala Thr Ile Leu Ser  
 450 455 460

Tyr Leu Thr Lys Ala Pro Leu Val Pro Pro Gly Thr Pro Val Val Asn  
 465 470 475 480

Ala Leu Ser Lys Gln Arg Ala Met Leu Glu Asn Ile Met Arg Ala Cys  
 485 490 495

Val Gly Leu Ala Pro Glu Asn Asn Met Ile Leu Glu Tyr Lys  
 500 505 510

(2) INFORMATION FOR SEQ ID NO:7:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 16 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: Synthetic oligonucleotide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

CGTAGGGGAC AGCAAG

16

(2) INFORMATION FOR SEQ ID NO:8:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 16 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: Synthetic oligonucleotide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

CGTAGGGGAC AGCAAT

16

## INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

<b>A.</b> The indications made below relate to the microorganism referred to in the description on page <u>6</u> , line s <u>31 &amp; 32</u>	
<b>B. IDENTIFICATION OF DEPOSIT</b> <span style="float: right;">Further deposits are identified on an additional sheet <input type="checkbox"/></span>	
Name of depositary institution <b>AMERICAN TYPE CULTURE COLLECTION</b>	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 US	
Date of deposit 17 April 1997 (17.04.97)	Accession Number 97988
<b>C. ADDITIONAL INDICATIONS</b> (leave blank if not applicable) <span style="float: right;">This information is continued on an additional sheet <input type="checkbox"/></span>	
In respect of those designations in which a European patent is sought, a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which the application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample. (Rule 28(4) EPC)	
<b>D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE</b> (if the indications are not for all designated States)	
<b>E. SEPARATE FURNISHING OF INDICATIONS</b> (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposit")	

<div style="text-align: center;"> <b>For receiving Office use only</b> </div> <div style="border: 1px solid black; padding: 5px;"> <input type="checkbox"/> This sheet was received with the international application  <b>G. Anton Smith</b>  <b>Paralegal Specialist</b>  <b>IAPD-PCT Operations</b>  <b>(703) 305-3747</b> </div> <div style="border: 1px solid black; padding: 5px; margin-top: 5px;"> Authorized officer </div>	<div style="text-align: center;"> <b>For International Bureau use only</b> </div> <div style="border: 1px solid black; padding: 5px;"> <input type="checkbox"/> This sheet was received by the International Bureau on: </div> <div style="border: 1px solid black; padding: 5px; margin-top: 5px;"> Authorized officer </div>
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## INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

<b>A. The indications made below relate to the microorganism referred to in the description</b> on page <u>6</u> , line s <u>31 &amp; 32</u>	
<b>B. IDENTIFICATION OF DEPOSIT</b> <span style="float: right;">Further deposits are identified on an additional sheet <input type="checkbox"/></span>	
Name of depositary institution AMERICAN TYPE CULTURE COLLECTION	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 US	
Date of deposit 04 April 1997 (04.04.97)	Accession Number 97971
<b>C. ADDITIONAL INDICATIONS</b> (leave blank if not applicable) <span style="float: right;">This information is continued on an additional sheet <input type="checkbox"/></span>	
In respect of those designations in which a European patent is sought, a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which the application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample. (Rule 28(4) EPC)	
<b>D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE</b> (if the indications are not for all designated States)	
<b>E. SEPARATE FURNISHING OF INDICATIONS</b> (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposit")	

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## INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

<b>A.</b> The indications made below relate to the microorganism referred to in the description on page <u>6</u> , line s <u>31 &amp; 32</u>	
<b>B. IDENTIFICATION OF DEPOSIT</b> <span style="float: right;">Further deposits are identified on an additional sheet <input type="checkbox"/></span>	
Name of depositary institution AMERICAN TYPE CULTURE COLLECTION	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 US	
Date of deposit 04 April 1997 (04.04.97)	Accession Number 97970
<b>C. ADDITIONAL INDICATIONS</b> (leave blank if not applicable) <span style="float: right;">This information is continued on an additional sheet <input type="checkbox"/></span>	
<p>In respect of those designations in which a European patent is sought, a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which the application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample. (Rule 28(4) EPC)</p>	
<b>D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE</b> (if the indications are not for all designated States)	
<b>E. SEPARATE FURNISHING OF INDICATIONS</b> (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposit")	

<p style="text-align: center;"><b>For receiving Office use only</b></p> <p><input type="checkbox"/> This sheet was received with the international application</p> <p style="text-align: center;"><b>Paralegal Specialist</b> <b>IAPD-PCT Operations</b></p> <p>Authorized officer <b>(703) 305-3747</b></p>	<p style="text-align: center;"><b>For International Bureau use only</b></p> <p><input type="checkbox"/> This sheet was received by the International Bureau on:</p> <p>Authorized officer</p>
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WHAT IS CLAIMED IS:

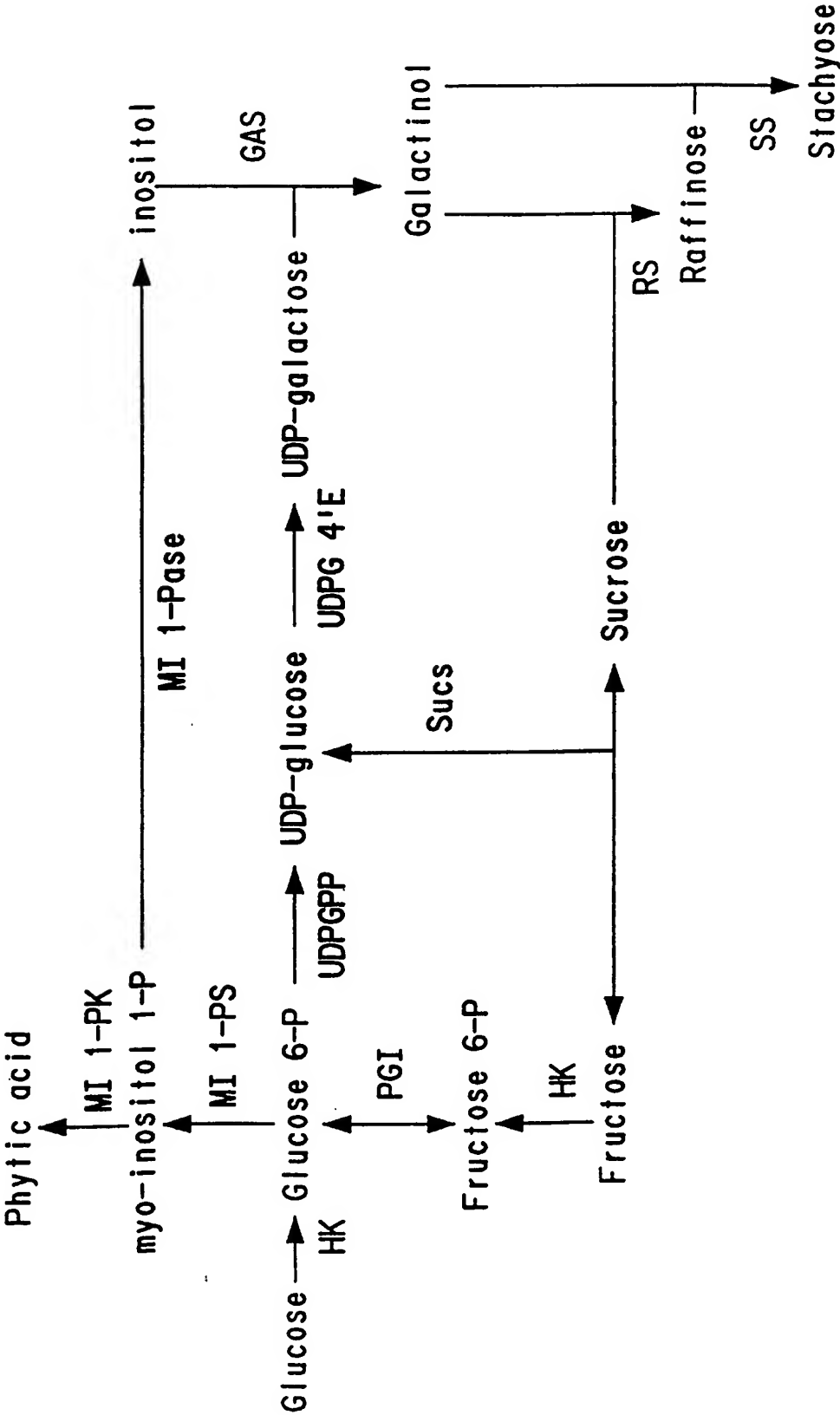
1. An isolated nucleic acid fragment encoding a soybean *myo*-inositol-1-phosphate synthase.
2. The nucleic acid fragment of Claim 1 wherein the amino acid sequence  
5 of the encoded soybean *myo*-inositol-1-phosphate synthase is substantially similar to the amino acid sequence set forth in SEQ ID NO:2.
3. A chimeric gene comprising the nucleic acid fragment of Claim 1 or the complement of the nucleic acid fragment of Claim 1, operably linked to suitable regulatory sequences, wherein expression of the chimeric gene results in a decrease in  
10 expression of a native gene encoding a soybean *myo*-inositol-1-phosphate synthase.
4. A chimeric gene comprising a subfragment of the nucleic acid fragment of Claim 1 or the complement of a subfragment of the nucleic acid fragment of Claim 1, operably linked to suitable regulatory sequences, wherein expression of the chimeric gene results in a decrease in expression of a native gene encoding a soybean  
15 *myo*-inositol-1-phosphate synthase.
5. An isolated nucleic acid fragment encoding a mutant *myo*-inositol 1-phosphate synthase having decreased capacity for the synthesis of *myo*-inositol-1-phosphate.
6. The nucleic acid fragment of Claim 5 wherein the amino acid sequence  
20 of the encoded mutant *myo*-inositol-1-phosphate synthase is substantially similar to the amino acid sequence set forth in SEQ ID NO:5.
7. A soybean plant homozygous for at least one gene encoding a mutant *myo*-inositol 1-phosphate synthase having decreased capacity for the synthesis of *myo*-inositol 1-phosphate, the gene conferring a heritable phenotype of (i) a seed phytic  
25 acid content of less than 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$ , provided that the plant is not LR33.
8. Seeds of the soybean plant of Claim 7.
9. A soybean plant comprising a member selected from the group  
30 consisting of:
  - (i) the chimeric gene of Claim 3;
  - (ii) the chimeric gene of Claim 4; and
  - (iii) the nucleic acid fragment of Claim 5;wherein the soybean plant has a heritable phenotype of (i) a seed phytic acid content  
35 less than 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$ , provided that the plant is not LR33.
10. Seeds of any of the soybean plants of Claim 9.

11. A method for making a soybean plant with a heritable phenotype of (i) a seed phytic acid content less than 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$ , the method comprising:
- 5 (a) crossing LR33 or any of the soybean plants of Claim 9 with an elite soybean plant; and
- (b) selecting progeny plants of the cross of step (a) that have a heritable phenotype of (i) a seed phytic acid content less than 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed
- 10 sucrose content of greater than 200  $\mu\text{mol/g}$ .
12. A soy protein product derived from seeds of a soybean plant homozygous for at least one gene encoding a mutant *myo*-inositol 1-phosphate synthase having decreased capacity for the synthesis of *myo*-inositol 1-phosphate, the gene conferring a heritable phenotype of (i) a seed phytic acid content less than
- 15 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$ .
13. A soy protein product derived from the processing of soybean seeds of Claim 8.
14. A soy protein product derived from the processing of soybean seeds of
- 20 Claim 10.
15. A method for producing a soy protein product derived from seeds of a soybean plant homozygous for at least one gene encoding a mutant *myo*-inositol 1-phosphate synthase having decreased capacity for the synthesis of *myo*-inositol 1-phosphate, the gene conferring a heritable phenotype of (i) a seed phytic acid content
- 25 less than 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$  comprising:
- (a) crossing an agronomically elite soybean plant with LR33 or any of the soybean plants of Claim 9;
- 30 (b) screening the seed of progeny plants obtained from step (a) for (i) a seed phytic acid content less than 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$ ; and
- (c) processing the seed selected in step (b) to obtain the desired
- 35 soybean protein product.
16. A method of using a soybean plant homozygous for at least one gene encoding a mutant *myo*-inositol 1-phosphate synthase having decreased capacity for the synthesis of *myo*-inositol 1-phosphate, the gene conferring a heritable phenotype of (i) a seed phytic acid content less than 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus

stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$  to produce progeny lines, the method comprising:

- (a) crossing a soybean plant comprising a mutant *myo*-inositol 1-phosphate synthase having decreased capacity for the synthesis of *myo*-inositol 1-phosphate with any soybean parent which does not comprise the mutation, to yield a F1 hybrid;
- (b) selfing the F1 hybrid for at least one generation; and
- (c) identifying the progeny of step (b) homozygous for at least one gene encoding a mutant *myo*-inositol 1-phosphate synthase having decreased capacity for the synthesis of *myo*-inositol 1-phosphate, the gene conferring a heritable phenotype of (i) a seed phytic acid content less than 17  $\mu\text{mol/g}$ , (ii) a seed content of raffinose plus stachyose combined of less than 14.5  $\mu\text{mol/g}$ , and (iii) a seed sucrose content of greater than 200  $\mu\text{mol/g}$ .

FIG. 1



# INTERNATIONAL SEARCH REPORT

Int. l. Application No  
PCT/US 98/06822

A. CLASSIFICATION OF SUBJECT MATTER  
IPC 6 C12N15/52 C12N15/82 C12N15/11 C12N9/90 A01H5/00

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
IPC 6 C12N A01H

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DEAN-JOHNSON, MARGARET ET AL: "Biosynthesis of inositol in yeast. Primary structure of myo-inositol-1-phosphate synthase (EC 5.5.1.4) and functional analysis of its structural gene, the INO1 locus" J. BIOL. CHEM. (1989), 264(2), 1274-83 CODEN: JBCHA3; ISSN: 0021-9258, XP002070761 * see esp. p.1277, p.1278 r. col. 1.21 - p.1279 r. col. 1.17 *	5,6
X	WO 93 07742 A (DU PONT) 29 April 1993 cited in the application see the whole document	12-14
A	WO 93 02196 A (DU PONT) 4 February 1993 see the whole document -/--	1-16

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

8 July 1998

Date of mailing of the international search report

20/07/1998

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# INTERNATIONAL SEARCH REPORT

International Application No  
PCT/US 98/06822

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>JOHNSON, MARGARET DEAN ET AL:  "1L-myo-Inositol 1-phosphate synthase from  Arabidopsis thaliana"  PLANT PHYSIOL. (1995), 107(2), 613-19  CODEN: PLPHAY;ISSN: 0032-0889, XP002070762  cited in the application  see the whole document  -----</p>	1-16

# INTERNATIONAL SEARCH REPORT

Information on patent family members

Inter. nat Application No

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